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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:28:19 ; Search time 222.28 Seconds
(without alignments)
347.126 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaagaagtgca 90

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 930621 segs, 428662619 residues

Word size: 0

Total number of hits satisfying chosen parameters: 989696

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

N.Geneseq_1101:*

1: /SIDS2/gcgdata/geneseq/geneseq/NA1980.DAT:*

2: /SIDS2/gcgdata/geneseq/geneseq/NA1981.DAT:*

3: /SIDS2/gcgdata/geneseq/geneseq/NA1982.DAT:*

4: /SIDS2/gcgdata/geneseq/geneseq/NA1983.DAT:*

5: /SIDS2/gcgdata/geneseq/geneseq/NA1984.DAT:*

6: /SIDS2/gcgdata/geneseq/geneseq/NA1985.DAT:*

7: /SIDS2/gcgdata/geneseq/geneseq/NA1986.DAT:*

8: /SIDS2/gcgdata/geneseq/geneseq/NA1987.DAT:*

9: /SIDS2/gcgdata/geneseq/geneseq/NA1988.DAT:*

10: /SIDS2/gcgdata/geneseq/geneseq/NA1989.DAT:*

11: /SIDS2/gcgdata/geneseq/geneseq/NA1990.DAT:*

12: /SIDS2/gcgdata/geneseq/geneseq/NA1991.DAT:*

13: /SIDS2/gcgdata/geneseq/geneseq/NA1992.DAT:*

14: /SIDS2/gcgdata/geneseq/geneseq/NA1993.DAT:*

15: /SIDS2/gcgdata/geneseq/geneseq/NA1994.DAT:*

16: /SIDS2/gcgdata/geneseq/geneseq/NA1995.DAT:*

17: /SIDS2/gcgdata/geneseq/geneseq/NA1996.DAT:*

18: /SIDS2/gcgdata/geneseq/geneseq/NA1997.DAT:*

19: /SIDS2/gcgdata/geneseq/geneseq/NA1998.DAT:*

20: /SIDS2/gcgdata/geneseq/geneseq/NA1999.DAT:*

21: /SIDS2/gcgdata/geneseq/geneseq/NA2000.DAT:*

22: /SIDS2/gcgdata/geneseq/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	14	15.6	20	AA205048	PCR primer used to
2	14	15.6	26	AAV07952	Helicobacter pylori
3	14	15.6	26	AAV07922	Helicobacter pylori
4	14	15.6	27	AAV07937	Helicobacter pylori
5	14	15.6	47	AA268842	Human map-related
6	13	14.4	17	AA092084	Human map-related
7	13	14.4	20	AA092084	Human map-related
8	13	14.4	23	AAH27610	PCR primer used to
9	13	14.4	24	AAH55939	Human SCNA PCR-SS
10	13	14.4	26	AA068537	B. thuringiensis 33
11	13	14.4	26	AA067312	Alzheimer's disease

12	13	14.4	26	AA067342	Alzheimer's disease
13	13	14.4	31	AAV07957	Helicobacter pylori
14	13	14.4	34	AAV15377	AZF A5F cosmid c10
15	13	14.4	35	AA061207	Coxsackie virus B
16	13	14.4	36	AA090560	Dissociation trans
17	13	14.4	38	AA134130	I3L promoter-HIV-
18	13	14.4	39	AA035630	HIV-2 env 3' fragm
19	13	14.4	39	AA035353	PCR primer HIV2B2
20	13	14.4	39	AA09138	Plasmid pomptc PC
21	13	14.4	40	AA048717	Insecticidal prote
22	13	14.4	40	AA261322	Primer 3A used to
23	13	14.4	40	AA261323	Primer 3B used to
24	13	14.4	41	AAV51009	Maize polymorphic
25	13	14.4	41	AAV51011	Maize polymorphic
26	13	14.4	50	AA025074	Human gene signatu
27	13	14.4	50	AA025184	Synthetic plasmid
28	13	14.4	50	AA025048	Synthetic plasmid
29	13	14.4	17	AA070046	Human fil1 VEGF re
30	12	13.3	17	AA070047	Human fil1 VEGF re
31	12	13.3	17	AA021453	Integrin alpha 6 s
32	12	13.3	17	AA021454	Integrin alpha 6 s
33	12	13.3	17	AA021455	Integrin alpha 6 s
34	12	13.3	17	AA030088	Hammerhead ribozym
35	12	13.3	17	AA030089	Hammerhead ribozym
36	12	13.3	17	AA030090	Hammerhead ribozym
37	12	13.3	17	AA030091	Hammerhead ribozym
38	12	13.3	18	AA089921	Human survivin DNA
39	12	13.3	19	AA062806	Env gene 5' primer
40	12	13.3	19	AA269625	Human biallelic ma
41	12	13.3	19	AA275126	Human biallelic ma
42	12	13.3	20	AA005907	HIV mRNA translati
43	12	13.3	20	AA080813	Staphylococcus aur
44	12	13.3	20	AAV72697	Corn kernel oil co
45	12	13.3	20	AA092327	PCR primer used to

ALIGNMENTS

RESULT 1	
AA205048	AA205048 standard; DNA; 20 BP.
XX	XX
XX	AA205048;
XX	07-OCT-1999 (first entry)
XX	XX
XX	XX
DE	PCR primer used to amplify an ORF of Chlamydia trachomatis.
XX	XX
KW	Vaccine, eye disease; conventional trachoma; nonendemic trachoma;
KW	paratrachoma; inclusion conjunctivitis; genital disease; perinephritis;
KW	nongonococcal urethritis; epididymitis; cervicitis; salpingitis; PCR primer;
KW	bartholinitis; pneumonia; venereal lymphogranulomatosis; ss.
XX	XX
OS	Synthetic.
OS	Chlamydia trachomatis.
XX	XX
PN	W09928475-A2.
PD	10-JUN-1999.
XX	XX
PF	27-NOV-1998; 98WO-IB01939.
XX	XX
PR	04-NOV-1998; 98US-0107077.
PR	28-NOV-1997; 97FR-0015041.
PR	17-DEC-1997; 97FR-0016034.
XX	XX
PA	(GEST) GENSET.
XX	XX
PI	Griffais R;
XX	XX
DR	WPI, 1999-371125/31.
XX	XX

PT Genome sequence of Chlamydia trachomatis
XX
PS Disclosure; Page 1738; 1755pp; English.
XX
CC PCR primers AAZ01426-206209 were used to amplify open reading frames
CC (ORFs) of the genome of Chlamydia trachomatis (see AAZ01425). These ORFs
CC encode polypeptides (see AAY36754-Y37949) which can be used as vaccines
CC against Chlamydia trachomatis. Antisense and ribozyme sequences
CC can also be used to control growth of the microorganism. Chlamydia
CC trachomatis is responsible for a large number of diseases, e.g. eye
CC diseases such as conventional trachoma, nongonococcal trachoma,
CC paratrachoma, and inclusion conjunctivitis; genital diseases such as
CC nongonococcal urethritis, epididymitis, cervicitis, salpingitis,
CC perihepatitis, Bartholinitis; pneumopathy in breast feeding infants;
CC and venereal lymphogranulomatosis. The polypeptides of the
CC invention may be of use in treating these diseases.
XX
SQ Sequence 20 BP; 5 A; 5 C; 5 G; 5 T; 0 other;

Query Match 15.6%; Score 14; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.4e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 54 tattgttgagca 67
|||||
Db 4 tattgttgagca 17

RESULT 2
AAV07952/C
ID AAV07952 standard; DNA; 26 BP.
XX
AC AAV07952;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHPO 1414 5' DNA primer.
XX
DE GHPO 1414: infection; gastritis; ulcer; vaccine; diagnosis;
XX therapy; PCR; primer; ss.
XX
KW Synthetic.
XX
OS Helicobacter pylori.
XX
OS WO9843479-A1.
XX
PN 08-OCT-1998.
XX
PD 31-MAR-1998; 98WO-US06421.
XX
PF 01-APR-1997; 97US-0834666.
XX
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PA (INNR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
PI WPI; 1998-568251/48.
XX
DR WPI; 1998-568251/48.
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
PS Claim 5; Page 145; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07954) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07921) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 1414.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter

CC Infections.
XX
SQ Sequence 26 BP; 14 A; 5 C; 4 G; 3 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 40 ttctcatgttct 53
|||||
Db 18 TTTTCATGTTTCT 5

RESULT 3
AAV07922/C
ID AAV07922 standard; DNA; 26 BP.
XX
AC AAV07922;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHPO 386 5' DNA primer.
XX
DE GHPO 386: infection; gastritis; ulcer; vaccine; diagnosis; therapy;
XX PCR; primer; ss.
XX
KW Synthetic.
XX
OS Helicobacter pylori.
XX
OS WO9843479-A1.
XX
PN 08-OCT-1998.
XX
PD 31-MAR-1998; 98WO-US06421.
XX
PF 01-APR-1997; 97US-0834666.
XX
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PA (INNR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
PI WPI; 1998-568251/48.
XX
DR WPI; 1998-568251/48.
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
PS Claim 5; Page 137; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07924) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV72001
CC) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 386. The
CC isolated polynucleotide, and encoded polypeptide, can be used to
CC develop vaccines for the treatment and prevention of Helicobacter
CC infections.
XX
SQ Sequence 26 BP; 15 A; 5 C; 4 G; 2 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 40 ttctcatgttct 53
|||||
Db 20 TTTTCATGTTTCT 7

RESULT 4
AAV07937/C
ID AAV07937 standard; DNA; 27 BP.
XX
AC AAV07937;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GHP0 896 5' DNA primer.
XX
KW GHP0 896; infection; gastritis; ulcer; vaccine; diagnosis;
KM therapy; PCR; primer; ss.
XX
OS Synthetic.
OS Helicobacter pylori.
XX
PM W09843479-A1.
XX
PD 08-OCT-1998.
XX
PF 31-MAR-1998; 98WO-US06421.
XX
PR 01-APR-1997; 97US-0834666;
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PA (IMMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
DR WPI; 1998-568251/48.
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
FT infections and gastroduodenal diseases
XX
PS Claim 5; Page 141; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07939) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07916) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73027) designated GHP0 896.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter
CC infections.
XX
SQ Sequence 27 BP; 14 A; 5 C; 4 G; 4 T; 0 other;

Query Match 15.6%; Score 14; DB 19; Length 27;
Best Local Similarity 100.0%; Pred. NO. 9.3e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 40 ttctcatgtttct 53
Db 18 TTTTCATCTTTCT 5
|||||
|||||

RESULT 5
AAZ68842/C
ID AAZ68842 standard; DNA; 47 BP.
XX
AC AAZ68842;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:3195.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KM genomic map; haplotype; phenotype; polymorphic base; genotyping;
KM haplotyping; hybridisation; identification; characterisation;
KM diagnosis; single nucleotide polymorphism; SNP; ds.
XX

OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,G)
FT /*tag= a
FT /standard_name= "single nucleotide polymorphism"
XX
PM W09954500-A2.
XX
PD 28-OCT-1999.
XX
PF 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
PI WPI; 2000-013267/01.
XX
DR Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome
XX
PS Claim 3; Page 912; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.
XX
SQ Sequence 47 BP; 13 A; 13 C; 6 G; 15 T; 0 other;

Query Match 15.6%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. NO. 9.1e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 aatgaagcaagt 88
Db 43 AATGAAGCAAGTG 30
|||||
|||||

RESULT 6
AAQ92084/C
ID AAQ92084 standard; CDNA; 17 BP.
XX
AC AAQ92084;
XX
DT 07-JAN-1996 (first entry)
XX
DE Renilla reniformis luciferase DNA probe-1.
XX
KW Luciferase; enzyme; bioluminescence; luminescence; label; DNA probe;
KM antibody; oligonucleotide; ss.
XX
OS Synthetic.
XX
PM US5418155-A.
XX

PD 23-MAY-1995.
 XX
 XX 29-DEC-1989; 89US-0458952.
 XX
 XX 29-DEC-1989; 89US-0458952.
 PR 20-AUG-1992; 92US-0933017.
 PR 17-JUN-1993; 93US-0079700.
 PR 14-DEC-1993; 93US-0167650.
 XX
 PA (UYGE-) UNIV GEORGIA RES FOUND INC.
 XX
 XX Cormier MJ, Lorenz WJ;
 PI
 XX WPI; 1995-199740/26.
 DR
 XX
 PT New recombinant Renilla luciferase polypeptide - used as a
 PT luminescent tag, partic in bio-luminescence assays and for the prodn
 PT of antibodies
 XX
 PS Disclosure; Fig. 4; 18pp; English.
 XX
 CC This 17-mer oligonucleotide DNA probe, along with probe-2 (AAQ92085)
 CC are used to screen an R. reniformis cDNA library to isolate cDNA
 CC encoding Renilla luciferase. The luciferase was then expressed
 CC using E. coli.
 XX
 SQ Sequence 17 BP; 6 A; 0 C; 2 G; 9 T; 0 other;

Query Match 14.4%; Score 13; DB 16; Length 17;
 Best Local Similarity 100.0%; Pred. No. 2.8e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattattt 16
 |||||
 Db 15 AAAAAATTTT 3

RESULT 7
 AAX95307
 ID AAX95307 standard; DNA; 20 BP.
 XX
 AC AAX95307;
 XX
 DT 13-SEP-1999 (first entry)
 XX
 DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.
 XX
 KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
 KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
 KW vaccine; neutralising epitope; PCR primer; ss.
 XX
 OS Synthetic.
 OS Chlamydia pneumoniae.
 OS
 PN WO9927105-A2.
 PN
 PD 03-JUN-1999.
 XX
 PF 20-NOV-1998; 98WO-IB01890.
 XX
 PR 04-NOV-1998; 98US-0107078.
 PR 21-NOV-1997; 97FR-0014673.
 XX
 PA (GEST) GENSET.
 XX
 PI Grifais R;
 XX
 DR WPI; 1999-357842/30.
 XX
 PT Genome sequence of Chlamydia pneumoniae
 PS Page 1737; Disclosure; 1912pp; English.

XX
 CC AAX91991-x97517 represent PCR primers used to amplify open reading
 CC frames and other nucleic acid sequences from the genome of
 CC Chlamydia pneumoniae (see AAX91990). C. pneumoniae causes respiratory
 CC disease such as pneumonia and bronchitis and is thought to be a
 CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent
 CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded
 CC by the open reading frames of the C. pneumoniae genome (see AAY34584-
 CC AAY35879) can be used in immunogenic compositions as vaccines. Vectors
 CC containing C. pneumoniae nucleotides sequences can also be used as
 CC immunogenic compositions, especially where the vector directs the
 CC expression of a neutralising epitope of C. pneumoniae.
 XX
 SQ Sequence 20 BP; 5 A; 5 C; 5 G; 5 T; 0 other;

Query Match 14.4%; Score 13; DB 20; Length 20;
 Best Local Similarity 100.0%; Pred. No. 2.8e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagtcg 89
 |||||
 Db 8 tgaagcaagtcg 20

RESULT 8
 AAH27610
 ID AAH27610 standard; DNA; 23 BP.
 XX
 AC AAH27610;
 XX
 DT 29-AUG-2001 (first entry)
 XX

DE Human lipoprotein 105 PCR primer 2.

XX
 KW Human; lipoprotein 105; cytosolic; anti-HIV; antiinflammatory;
 KW immunomodulatory; cytochrome c structural domain; cancer;
 KW haemopathy; human immunodeficiency virus; HIV; immunological disease;
 KW inflammation; PCR primer; ss.
 XX

OS Homo sapiens.

XX
 PN WO200140483-A1.
 PN
 PD 07-JUN-2001.
 XX

PF 27-NOV-2000; 2000WO-CN00503.

XX
 PR 29-NOV-1999; 99CN-0124136.

XX
 PA (BIOR-) BIOROAD GENE DEV LTD SHANGHAI.

XX
 PI Mao Y, Xie Y;
 XX

DR WPI; 2001-374839/39.
 DR

PT Human lipoprotein 105 containing cytochrome c structural domain and
 PT encoded polynucleotide, applicable in diagnosis and treatment of
 PT malignant tumor, hemopathy, HIV infection, immunological diseases and
 PT various inflammation
 XX
 PS Example 2; Page 12; 40pp; Chinese.
 XX

CC The invention relates to an isolated polypeptide of human lipoprotein
 CC 105 containing a cytochrome c structural domain. The polypeptide
 CC comprises a 951 amino acid sequence given in the specification, or its
 CC fragment, analogue or derivative. The polypeptide and encoded
 CC polynucleotide are useful in the diagnosis and treatment of malignant
 CC tumours, haemopathy, HIV infection, immunological diseases and various
 CC inflammatory diseases. The present sequence is a primer which was
 CC used to isolate the polynucleotide encoding the polypeptide of the
 CC invention.

Sequence 23 BP; 5 A; 1 C; 2 G; 15 T; 0 other;

Query Match 14.4%: Score 13; DB 22; Length 23;

Best Local Similarity 100.0%: Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

33 ttattatgtttca 45
|||||
1 ttattatgtttca 13

RESULT 9

AAH55939

ID AAH55939 standard; DNA; 24 BP.

AAH55939;

04-SEP-2001 (first entry)

Human SCN1A PCR-SSCP PCR primer SEQ ID NO:183.

Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification; diagnosis; mutation; chromosome 2q23-q31; neurological disorder; anticonvulsant; neuroprotective; PCR primer; ss.

Homo sapiens.
Synthetic.

WO200138564-A2.

31-MAY-2001.

24-NOV-2000; 2000WO-CA01404.

26-NOV-1999; 99US-0167623.

(UYMC-) UNIV MCGILL.

Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;

WPI; 2001-355945/37.

Determining a predisposition to epilepsy and/or development of epilepsy comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA variant, equivalent, or mutation which shows a linkage disequilibrium -

Example 3; Fig 2; 268pp; English.

The present invention describes a method (M1) of determining an individual's predisposition to epilepsy and/or development of epilepsy, as well as predicting the individual's response to medication. The method comprises determining the genotype of at least one gene selected from SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium channel genes located on chromosome 2. The idiopathic generalised epilepsy (IGE) gene is more specifically localised on chromosome 2q23-q31. Compounds identified as modulators of the biological activity of SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy or other neurological disorders. They have anticonvulsant and neuroprotective activities. AAH55763 to AAH56164 and AAH99674 to AAH99679 represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers, oligonucleotides and proteins given in the exemplification of the present invention.

Sequence 24 BP; 2 A; 5 C; 3 G; 14 T; 0 other;

Query Match 14.4%: Score 13; DB 22; Length 24;

Best Local Similarity 100.0%: Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 34 ttatgtttcat 46
|||||
Db 8 ttatgtttcat 20

RESULT 10

AAO68537/C

ID AAO68537 standard; cDNA; 26 BP.

AAO68537;

13-FEB-1995 (first entry)

B.thuringiensis 33kD delta-endotoxin N-terminal probe.

Insecticidal protein; delta-endotoxin; crystal; Coleoptera; Lepidoptera; Bacillus thuringiensis; ss.

Synthetic.

WO9413785-A.

23-JUN-1994.

13-DEC-1993; 93WO-US12144.

15-DEC-1992; 92US-0991073.

(NOVO) NOVO-NORDISK ENTOTEC INC.

Adams LF, Liu C, Lufurrow PA, Thomas MD;

WPI; 1994-217865/26.

New Bacillus thuringiensis strains - which produce new delta-endotoxin cpds used for the control of Lepidopteran and Coleopteran insect pests.

Example 9; Page 26; 47pp; English.

The N-terminal sequence of a 33kD delta-endotoxin isolated from B.thuringiensis EMCC0075 (NRRL B-21019) or EMCC0076 (NRRL B-21020) was determined (see AAR59764). Based on this sequence a 26mer oligonucleotide was designed for use as a probe (AAO68537) for cloning the delta-endotoxin gene.

Sequence 26 BP; 13 A; 2 C; 2 G; 9 T; 0 other;

Query Match 14.4%: Score 13; DB 15; Length 26;
Best Local Similarity 100.0%: Pred. No. 2.8e+03; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 30 aattttatgttt 42
|||||
Db 16 AATTTTATGTTT 4

RESULT 11

AAC67312

ID AAC67312 standard; DNA; 26 BP.

AAC67312;

14-FEB-2001 (first entry)

Alzheimer's disease-linked mitochondrial SNP PCR primer #12.

Human: mitochondrial genome; single nucleotide polymorphism; SNP; Alzheimer's disease; mtDNA; PCR primer; ss.

Homo sapiens.

PN WO200063441-A2.
XX
PD 26-OCT-2000.
XX
PF 19-APR-2000; 2000WO-US10906.
XX
PR 20-APR-1999; 99US-0130447.
PR 22-OCT-1999; 99US-0160901.
XX
PA (MITO-) MITOKOR.
PI Herrnstadt C, Davis RE;
XX
DR WPI; 2000-672748/65.
XX
PT Diagnosing a subject at the risk for or having Alzheimer's disease
PT comprises determining at least one single nucleotide polymorphism in
PT mitochondrial DNA associated with the disease in the sample from the
PT subject -
XX
PS Example 2; Page 34; 89pp; English.
XX
CC The present invention describes a novel method for determining the risk
CC of or diagnosing Alzheimer's disease using single nucleotide
CC polymorphisms (SNPs) present in an individual's mitochondrial DNA
CC (mtDNA). In addition, the SNPs identified can be used to identify agents
CC suitable for use in treating Alzheimer's disease. Sequences
CC AAC67301-C67610 are PCR primers used to demonstrate the method of the
CC invention.
XX
SQ Sequence 26 BP; 8 A; 2 C; 5 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 37 atgtttcatgtt 49
|||||
Db 7 atgtttcatgtt 19

RESULT 12
AAC67342
ID AAC67342 standard; DNA; 26 BP.
XX
AC AAC67342;
XX
DT 14-FEB-2001 (first entry)
XX
DE Alzheimer's disease-linked mitochondrial SNP PCR primer #42.
XX
KW Human; mitochondrial genome; single nucleotide polymorphism; SNP.
KW Alzheimer's disease; mtDNA; PCR primer; ss.
XX
OS Homo sapiens.
XX
PN WO200063441-A2.
XX
PD 26-OCT-2000.
XX
PF 19-APR-2000; 2000WO-US10906.
XX
PR 20-APR-1999; 99US-0130447.
PR 22-OCT-1999; 99US-0160901.
XX
PA (MITO-) MITOKOR.
PI Herrnstadt C, Davis RE;
XX
DR WPI; 2000-672748/65.
XX
PT Diagnosing a subject at the risk for or having Alzheimer's disease

PT comprises determining at least one single nucleotide polymorphism in
PT mitochondrial DNA associated with the disease in the sample from the
PT subject -
XX
PS Example 4; Page 38; 89pp; English.
XX
CC The present invention describes a novel method for determining the risk
CC of or diagnosing Alzheimer's disease using single nucleotide
CC polymorphisms (SNPs) present in an individual's mitochondrial DNA
CC (mtDNA). In addition, the SNPs identified can be used to identify agents
CC suitable for use in treating Alzheimer's disease. Sequences
CC AAC67301-C67610 are PCR primers used to demonstrate the method of the
CC invention.
XX
SQ Sequence 26 BP; 8 A; 2 C; 5 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 37 atgtttcatgtt 49
|||||
Db 7 atgtttcatgtt 19

RESULT 13
AAV07957/G
ID AAV07957 standard; DNA; 31 BP.
XX
AC AAV07957;
XX
DT 02-FEB-1999 (first entry)
XX
DE Helicobacter pylori polypeptide GPO 386 5' DNA primer.
XX
DE GPO 386; infection; gastritis; ulcer; vaccine; diagnosis; therapy;
KW PCR; primer; ss.
KW
OS Synthetic.
OS Helicobacter pylori.
XX
PN WO9843479-A1.
XX
PD 08-OCT-1998.
XX
PF 31-MAR-1998; 98WO-US06421.
XX
PR 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HOMA-) HUMAN GENOME SCI INC.
PA (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX
PI Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX
DR WPI; 1998-568251/48.
XX
PT New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
PS Example 3.B; Page 63; 184pp; English.
XX
CC This 5' primer was used with a 3' primer (see AAV07958) in the PCR
CC amplification of Helicobacter pylori strain ORV2001 genomic DNA in
CC order to obtain DNA (see AAV72001
CC) encoding a 76 kDa polypeptide (see
CC AAV73022) designated GPO 386. The primer pair includes a 5' clamp
CC and BamHI and XhoI restriction enzyme recognition sequences for
CC cloning purposes. The PCR product was ligated into vector pET28a,
CC and recombinant polypeptide was expressed as a histidine-tagged
CC fusion protein in E. coli host cells. The polypeptide can be used

CC to develop vaccines for the treatment and prevention of Helicobacter
 CC infections.

XX Sequence 31 BP; 14 A; 4 C; 6 G; 7 T; 0 other;

XX SQ

Query Match 14.4%; Score 13; DB 19; Length 31;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 41 ttcatgttttct 53
 |||||
 DB 31 TTTCATGTTTCT 19

RESULT 14

AA15377
 ID AA15377 standard; DNA; 34 BP.

XX AC AA15377;

XX DT 20-MAR-1996 (first entry)

XX DE AZF A5F cosmid clone exon 9/intron 9 boundary.

XX KW Azospermia factor; AZF; male infertility; YRRM gene;

XX KM Y-chromosome; ss.

XX OS Homo sapiens.

XX Key Location/Qualifiers

XX FT exon 1..17

XX FT intron /tag= a /number= 9

XX FT /note= "Represents bases 1093-1109 of human YRRM gene"

XX FT /tag= b /number= 9

XX FT /note= "Constitutes 5' end of approx. 450 bp intron 9"

XX PN W09511300-A2.

XX PD 27-APR-1995.

XX XX 24-OCT-1994; 94WO-GB02344.

XX PR 07-JUL-1994; 94GB-0013760.

XX PR 22-OCT-1993; 93GB-0021857.

XX PA (MED1-) MEDICAL RES COUNCIL.

XX PI Chandley AC, Cooke HJ, Hargreave TB, Kun M, Sharkey AM;

XX DR WPI: 1995-170221/22.

XX PS Nucleic acid encoding the human azospermia factor, and probes and

XX PT antibodies specific for the sequence and encoded polypeptide - may

XX PT be used in the clinical diagnosis of male infertility

XX PS Disclosure: Fig 3; 40pp; English.

XX CC The intron/exon boundaries of a human YRRM gene encoding

XX CC azospermia factor were detd. by comparison of the sequences

XX CC found in cosmid A5F DNA obtd. from a Y-chromosome specific cosmid

XX CC library and the cDNA clone MK5 (AA087655) obtd. from an adult human

XX CC testis library.

XX SQ Sequence 34 BP; 9 A; 3 C; 7 G; 15 T; 0 other;

OY 24 tactgtaatttt 36
 |||||
 DB 19 tactgtaatttt 31

RESULT 15

AA61207/c
 ID AA61207 standard; DNA; 35 BP.

XX AC AA61207;

XX DT 25-MAY-2001 (first entry)

XX DE Cocksackie virus B serotype B3 associated PCR primer SEQ ID 4.

XX KW Gene therapy; Infectious virion; cardiactive; cardiac muscle; trophic;

XX KW cardiac disease; cardiac myocyte; PCR primer; ss.

XX OS Cocksackievirus.

XX PN DE19939095-A1.

XX PD 22-FEB-2001.

XX PR 18-AUG-1999; 99DE-1039095.

XX PR 18-AUG-1999; 99DE-1039095.

XX PA (UYTU-) UNIV TUEBINGEN EBERHARD-KARLS.

XX PI Kuepper J, Meyer R, Meyer-Ficca M, Kandolf R;

XX DR WPI: 2001-227548/24.

XX PT Recombinant RNA comprising heterologous gene in Cocksackie viral genome,

XX PT useful in gene therapy, specifically for targeting of cardiac myocytes

XX PT Example 3; Column 10; 12pp; German.

XX PS This invention describes a novel recombinant RNA molecule (I), at least

XX CC partly translatable in a target cell which comprises: (a) the

XX CC non-infectious genome (A) of Group B Cocksackie virus (CVB), particularly

XX CC serotype B3; and (b) at least one foreign gene (II) that can be developed

XX CC for a selected function in the target cell, e.g. for gene therapy. The

XX CC invention also describes (1) a recombinant infectious virions (V),

XX CC derived from (A) and containing (I); (2) a plasmid vector containing the

XX CC DNA sequence (III) for (1), under control of a promoter; (3) a helper

XX CC construct for complementing the coding sequence exchanged by (1); (4)

XX CC producing (V); (5) producing the plasmids of (2); (6) producing helper

XX CC constructs of (3); (7) a kit containing the vector of (2) or the helper

XX CC construct of (3); (8) a DNA molecule (IV) containing at least one coding

XX CC sequence for (I); (9) a kit containing (IV); (10) a kit for performing

XX CC methods (5) or (6); (11) a DNA construct that encodes (I) and can persist

XX CC (12) and is transcribed in target cells, but is preferably not replicable;

XX CC (13) a recombinant virus (RV), particularly adeno or retro, that encodes

XX CC (I) and is expressed after infection into a target cell to produce a

XX CC cytoplasmic replicon that is continuously replenished; (13) producing

XX CC recombinant DNA viruses or virions having a DNA genome that lacks a

XX CC specific gene function, in which this function is provided from a

XX CC recombinant vector system with a RNA genome. The products of the

XX CC invention have cardioactive activity and can be used for gene therapy.

XX CC (I) is used to produce gene therapy vectors, particularly plasmids or

XX CC virions, for diagnosis, prevention or treatment of cardiac disease, either

XX CC muscle, for diagnosis, prevention or treatment of cardiac disease, either

XX CC congenital or acquired. (I) Are also used to complement vectors that lack

XX CC particular gene sequences, particularly vectors derived from DNA viruses.

XX CC Vectors based on (I) transfer genes to cardiac myocytes without

XX CC immunological or other side effects. The RNA genome can replicate,

XX CC providing efficient gene transfer and long-term expression of the

XX CC therapeutic gene. CBV is naturally tropic for heart muscle and since it

XX CC does not produce DNA during its life cycle, overcomes the danger that

CC foreign genes will become integrated in the target cell genome. By using
 CC (II) to replace part of the viral coding region, large (II) sequences may
 CC be accommodated. (I) is easily packaged in CVB capsid proteins.
 XX
 SO Sequence 35 BP; 14 A; 2 C; 7 G; 12 T; 0 other;

Query Match 14.4%; Score 13; DB 22; Length 35;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 aaattattcaaa 20
 |||
 Db 27 AAATATTTCAAA 15

RESULT 16

AAF90560
 ID AAF90560 standard; DNA; 36 BP.

AC AAF90560;

DT 22-AUG-2001 (first entry)

DE Dissociation transposon 3' flanking nested primer 3A.

KW ET1158 gene; GT6839 gene; ET5262 gene; herbicide; screening;

KW herbicide tolerance; transgenic plant; crop protection;

KW Dissociation; transposon; maize; PCR primer; ss.

OS Zea mays.

PN MO200144277-A2.

PD 21-JUN-2001.

PE 14-DEC-2000; 2000WO-EP12748.

PR 16-DEC-1999; 99US-0465040.

PI (SYGN) SYNGENTA PARTICIPATIONS AG.

DR Wegrich Glover L, Budziszewski GJ, Levin JZ, Zhou Q;

XX MPI; 2001-398122/42.

XX New herbicide target genes encoding proteins having ET1158, GT6839 or
 PT ET5262 activity, for identifying an inhibitor of protein activity -

PS Example 1; Page 39; 67pp; English.

CC The present sequence is that of primer 3A, which is 1 of a set of
 CC 3 nested primers (see also AAF90561 and AAF90562) homologous to
 CC the 3' DS3 region of Dissociation transposon. Arbitrary degenerate
 CC primers (see AAF90551-56) were used to prime Arabidopsis thaliana
 CC genomic DNA flanking the site of a DS transposon insertion. The
 CC degenerate primers were used in combination with 2 sets of 3,
 CC nested, transposon-specific primers homologous to the DS3 region or
 CC DS5 region (see AAF90557-59) of the DS element, which lie at the
 CC outermost ends of the transposon. Low- and high-stringency PCR
 CC amplifications were performed using the TAIL-PCR protocol. DNA
 CC fragments were produced which corresponded to the genomic DNA that
 CC was directly adjacent to the transposon insertion. Sequence
 CC analysis of PCR products from tagged seedling lethal lines ET1158,
 CC GT6839 and ET5262 identified 3 novel genes (see AAF90548-50) each
 CC of which was essential for Arabidopsis seedling growth and
 CC development. The essentiality of the genes provides a means of
 CC discovering new herbicides. Screening assays for identifying
 CC inhibitors that are potential herbicides are provided. The
 CC invention is also applied to the development of herbicide tolerant
 CC plants, and plant tissues, seeds and cells.

XX Sequence 36 BP; 6 A; 4 C; 11 G; 15 T; 0 other;

Query Match 14.4%; Score 13; DB 22; Length 36;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 36 tatgttttcattgt 48
 |||
 Db 21 tatgttttcattgt 33

RESULT 17

AAF34130/c
 ID AAT34130 standard; cDNA; 38 BP.

AC AAT34130;

DT 21-OCT-1996 (first entry)

DE I3L promoter-HTLV-I fusion primer MW116.

KW Avipox virus; poxvirus; vector; vaccine; attenuation; HTLV-I

KW human lymphotropic virus; safety; ALVAC; NYVAC; primer; PCR;

KW polymerase chain reaction; vaccinia virus; retrovirus; ss.

OS Synthetic.

PN WO9621727-A1.

PD 18-JUL-1996.

PE 16-JAN-1996; 96WO-US00547.

PR 13-JAN-1995; 95US-0372664.

PI (VIRO-) VIROGENETICS CORP.

DR Franchini G, Gallo RC, Paolletti E, Tartaglia J;

XX MPI; 1996-342282/34.

XX Attenuated recombinant pox-viruses expressing HTLV antigens - for
 PT safe vaccination against HTLV infection, and for therapy and
 PT diagnosis

PS Example 12; Page 96; 165pp; English.

CC Primers MW093 (AAT34127) and MW110 (AAT34128) were used to generate
 CC a 100 bp PCR fragment, PCR-HTLV18, contg. the I3L promoter fused to
 CC the 5'-end of the HTLV-I envelope gene, using PMM102 as template.
 CC Primers MW113 (AAT34129) and MW116 (AAT34130) were used to generate a
 CC 1,500 bp PCR fragment, PCR-HTLV21, contg. the 3' end of the I3L
 CC promoter fused to the entire HTLV-I envelope gene, using p17-SST as
 CC template. MW093 and MW116 were then used as to amplify PCR-HTLV18
 CC and PCR-HTLV21. The I3L-promoted envelope gene was cloned between
 CC canarypox virus C5 flanking arms and also between vaccinia HA
 CC flanking arms, and a NYVAC recombinant expressing HTLV-I env was
 CC generated. This effectively primed protective immune responses
 CC against retrovirus challenge.

XX Sequence 38 BP; 15 A; 6 C; 8 G; 9 T; 0 other;

Query Match 14.4%; Score 13; DB 17; Length 38;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
 |||
 Db 25 CTTGTAATTTTAT 13

RESULT 18

AAQ35630/C
ID AAQ35630 standard; DNA; 39 BP.
XX
AC AAQ35630;
XX
XX 24-FEB-1993 (first entry)
XX
DE HIV-2 env 3' fragment primer HIV2B2.
XX
XX NYVAC; recombinant; HIV-2; Copenhagen vaccine; vaccinia virus;
KM virulence factors; deletion loci; recipient loci; env; amplify; PCR;
KM polymerase chain reaction; gp120; transfection; Vero cells; envelope;
KM cell surface; ss.
XX
OS Synthetic.
XX
XX WO9215672-A.
XX
PD 17-SEP-1992.
XX
PF 09-MAR-1992; 92WO-US01906.
XX
XX 07-MAR-1991; 91US-0666056.
PR 11-JUN-1991; 91US-0713967.
PR 06-MAR-1992; 92US-0847951.
XX
PA (VIRO-) VIROGENETICS CORP.
XX
XX Cox WI, De Taisne C, Francis J, Gettig RR, Johnson GP;
PI Limbach KJ, Norton EK, Paoletti E, Perkus ME, Pincus SE;
PI Riviere M, Tartaglia J, Taylor J;
XX
XX WPI, 1992-331718/40.
XX
PT Vaccine comprises recombinant, attenuated pox-virus - use for
PT vaccinating against viral infections such as rabies, hepatitis B,
PT HIV, HSV, EBV, CMV, mumps etc.
XX
PS Disclosure; Page 156; 455pp; English.
XX
XX The sequences given in AAQ35624-32 were used in the construction of
CC NYVAC recombinants expressing HIV-2 gene products. NYVAC is a
CC Copenhagen vaccine strain of vaccinia virus which has been modified by
CC deletion of six non-essential regions of the genome encoding known or
CC potential virulence factors. The deletion loci were engineered as
CC recipient loci for the insertion of foreign genes. The HIV-2 env
CC sequence was isolated in two portions by polymerase chain reaction
CC (PCR). These fragments were then ligated also by PCR. The HIV-2
CC gp120 gene was also isolated by PCR. These HIV-2 genes were
CC transfected into NYVAC which could then be cultured in Vero cells.
CC The envelope proteins were found to be present on the cell surface of
CC cells transformed with the recombinant NYVAC. See AAQ35501-864.
XX
SQ Sequence 39 BP; 15 A; 6 C; 7 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 13; Length 39;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
|||||
DB 25 CTGTAATTTTAT 13

RESULT 19
AAQ35353/C
ID AAQ35353 standard; DNA; 39 BP.
XX
AC AAQ35353;
XX
XX 18-MAY-1993 (first entry)
XX

DE PCR primer HIV2B2.
XX
XX Human immunodeficiency virus; amplification; env; ISSY strain; ss.
XX
XX Synthetic.
XX
XX WO9222641-A.
XX
XX 23-DEC-1992.
XX
XX 12-JUN-1992; 92WO-US05107.
PF 14-JUN-1991; 91US-0715921.
PR 11-JUN-1992; 92US-0897382.
XX
XX (VIRO-) VIROGENETICS CORP.
XX
XX Cox WI, Paoletti E, Tartaglia J;
XX
XX WPI, 1993-018128/02.
XX
XX Modified recombinant virus with inactivated non-essential genetic
PT functions - comprises e.g. vaccinia or avipox virus, used as HIV
PT vaccine
XX
XX Example 4; Page 52; 159pp; English.
XX
XX The 3' portion of the HIV-2 env gene was derived by PCR. In this
CC reaction a 270 bp fragment was amplified using oligonucleotides
CC HIV2B1 and HIV2B2 using P1SSY-KPN as template. The prod. was
CC digested with BamHI and XbaI to yield a 150 bp fragment which was
CC engineered to contain a T5NT sequence known to be recognised as
CC vaccinia virus early transcription termination signal, following
CC the termination codon TAA.
XX
XX See also AAQ35328-437.
XX
SQ Sequence 39 BP; 15 A; 6 C; 7 G; 11 T; 0 other;

Query Match 14.4%; Score 13; DB 14; Length 39;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
|||||
DB 25 CTGTAATTTTAT 13

RESULT 20
AAA99138
ID AAA99138 standard; DNA; 39 BP.
XX
XX AAA99138;
XX
XX 19-JAN-2001 (first entry)
XX
XX Plasmid pOMPTTC PCR primer SEQ ID NO:15.
DE OMP1 protease; cleavage; fusion protein; membrane protease;
XX natriferic; Escherichia coli; PCR primer; ss.
KM natriferic; Escherichia coli; PCR primer; ss.
XX
XX Escherichia coli.
OS Synthetic.
XX
XX WO200052193-A1.
XX
XX 08-SEP-2000.
XX
XX 03-MAR-2000; 2000WO-JP01309.
PF 04-MAR-1999; 99JP-0057731.
XX
XX (SUNR) SUNTORY LTD.
PA

```

XX Okuno K, Yabuta M, Ohsuye K;
PI WPI: 2000-579291/54.
XX
XX Controlled cleavage of peptides by OmpT protease by amino acid
PT substitution for ensuring cleavage only at desired site in fission of
XX fusion proteins
XX
XX Example 2; Fig 9; 144pp; Japanese.
XX
XX The present invention describes a method for regulating the cleavage
CC sites of polypeptides by OmpT protease by preventing cleavage at
CC unwanted sites by converting the amino acid residue at position +1 to
CC the site to a specifically defined amino acid (where the residue at
CC position -1 to the site is Lys or Arg), and/or converting the residue
CC at position -4 and/or -6 to a specifically defined amino acid. Also
CC described is a method for the fission of a fusion protein to give a
CC desired polypeptide by cleavage with OmpT protease, where the fusion
CC protein has a linker peptide inserted between the desired polypeptide
CC and the other part of the fusion protein. The fusion protein may be
CC prepared by expression of DNA encoding in a suitable host cell such as
CC Escherichia coli. OmpT protease is a membrane protease of Escherichia
CC coli which cleaves peptide chains at a two-residue sites in which both
CC residues are basic amino acids such as arginine or lysine. The methods
CC can be used for the efficient preparation of undegraded desired
CC polypeptides such as natural peptide by OmpT protease cleavage after
CC recombinant expression as a fusion protein. AA99127 to AA99177 and
CC AAB3946 to AAB24018 represent sequences used in the exemplification of
CC the present invention.
XX
XX Sequence 39 BP; 8 A; 9 C; 4 G; 18 T; 0 other:
SQ

```

Query Match 14.4%; Score 13; DB 21; Length 39;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

OY 26 cgtatattttat 38
DB 15 cgtatattttat 27

```

RESULT 21
 AA048717/c
 ID AA048717 standard; DNA; 40 BP.
 XX
 XX AA048717;
 AC
 XX
 XX 25-MAR-1994 (first entry)
 DT
 XX
 XX Insecticidal protein gene 3' primer.
 DE
 XX Caulobacter; plasmid; insecticidal protein; *Bacillus thuringiensis*;
 KW *Bacillus sphaericus*; larva; mosquito; *Culex*; *Anopheles*; *Psorophora*;
 KW *Mansonia*; *Aedes*; ss.
 XX
 XX Synthetic.
 OS
 XX
 XX JP05211866-A.
 PN
 XX
 XX 24-AUG-1993.
 PD
 XX
 XX 05-JUN-1991; 91JP-0160963.
 PF
 XX
 XX 06-JUN-1990; 90JP-0148444.
 PR
 XX (SILM-) SILMARAN SO TANABAL.
 PA
 XX WPI: 1993-298916/38.
 DR
 XX
 XX Expression of insecticidal protein - by transforming *Caulobacter*
 PT with plasmid contg. gene coding for insecticidal protein

```

XX Disclosure; Page 24-25; 27pp; Japanese.
PS
XX Two primers (AA048716-17) are described for the isolation of
CC Bacillus thuringiensis israelensis DNA. The sequence of one
CC 27-mer primer (AA048716), however is missing from the specification.
CC Caulobacter transformed with a plasmid contg. a gene encoding
CC insecticidal protein derived from Bacillus thuringiensis or
CC Bacillus sphaericus will proliferate in aq. environment.
CC They may be consumed by larvae of mosquitoes and are lethal to
CC Culex, Anopheles, Psorophora, Mansonia and Aedes.
XX
XX Sequence 40 BP; 19 A; 6 C; 6 G; 9 T; 0 other:
SQ

```

Query Match 14.4%; Score 13; DB 14; Length 40;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

OY 32 tttrtggtttc 44
DB 15 tttrtggtttc 3

```

RESULT 22
 AA261322/c
 ID AA261322 standard; DNA; 40 BP.
 XX
 XX AA261322;
 AC
 XX
 XX 19-JUN-2000 (first entry)
 DT
 XX
 XX Primer 3A used to construct a synthetic bar gene.
 DE
 XX bar gene; phosphinothricin acetyltransferase; phosphinothricin toxicity;
 KW protein expression; vaccine; haemoglobin; enzyme; primer; ss.
 KW
 OS Synthetic.
 OS Streptomyces hygroscopicus.
 OS
 XX
 XX WO200007431-A1.
 PN
 XX
 XX 17-FEB-2000.
 PD
 XX
 XX 03-AUG-1999; 99WO-US17806.
 PF
 XX
 XX 03-AUG-1998; 980S-0095163.
 PR
 XX 03-AUG-1998; 980S-0095167.
 PR
 XX 15-DEC-1998; 980S-0112257.
 PR
 XX 29-APR-1999; 990S-0131611.
 PR
 XX 11-JUN-1999; 990S-0138764.
 XX
 XX (RUTP) UNIV RUTGERS STATE NEW JERSEY.
 PA
 XX
 XX Maliga P, Kuroda H, Khan MS;
 PI
 XX
 XX WPI: 2000-205525/18.
 DR
 XX
 XX New recombinant DNA constructs, for expressing high levels of
 PT heterologous protein in plasmids of higher plants, includes promoter, a
 PT leader sequence and a downstream box element -
 XX
 XX Example 7; Page 69; 164pp; English.
 PS
 XX
 XX Primers AA261218-45 were used to construct a synthetic *Streptomyces*
 CC hygroscopicus bar gene. The bar gene encodes phosphinothricin
 CC acetyltransferase, which provides protection from phosphinothricin
 CC toxicity. The synthetic gene has improved containment and enhanced
 CC expression in plant plasmids, and is used to produce recombinant DNA
 CC constructs of the invention. The specification describes recombinant DNA
 CC constructs for expressing heterologous proteins (e.g. bar gene product)
 CC in the plasmids of higher plants. The DNA constructs comprise a 5'
 CC regulatory region which includes a promoter element, a leader sequence

CC and a downstream box element operably linked to a coding region of the
CC heterologous protein. The chimeric regulatory region enhances
CC translational efficiency of an mRNA molecule encoded by the DNA
CC construct. The DNA constructs are used for producing transformed
CC monocot and dicot plants having high levels of heterologous protein
CC expression. They can be used to drive expression of proteins with
CC agronomic, industrial or pharmaceutical importance, including production
CC of vaccines, healthcare products like human haemoglobin, industrial or
CC household enzymes. Plants which can be transformed with the constructs
CC of the invention include maize, millet, sorghum, sugar cane, rice,
CC wheat, barley, oat, rye or turf grass.
XX
SQ Sequence 40 BP; 19 A; 6 C; 5 G; 10 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 40;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 18 aaagtctactgta 30
|||||
Db 33 AAAGTTTACTGTA 21

RESULT 23
AA61323
ID AA61323 standard; DNA; 40 BP.
XX
AC AA61323;
XX
DT 19-JUN-2000 (first entry)
XX
DE Primer 3B used to construct a synthetic bar gene.
XX
KW bar gene; phosphinothricin acetyltransferase; phosphinothricin toxicity;
KW protein expression; vaccine; haemoglobin; enzyme; primer; ss.
XX
OS Synthetic.
OS Streptomyces hygroscopicus.
XX
PN WO200007431-A1.
XX
PD 17-FEB-2000.
XX
PF 03-AUG-1999; 99WO-US17806.
XX
PR 03-AUG-1998; 98US-0095163.
PR 03-AUG-1998; 98US-0095167.
PR 15-DEC-1998; 98US-0112257.
PR 29-APR-1999; 99US-0131611.
PR 11-JUN-1999; 99US-0138764.
XX
PA (RUTE) UNIV RUTGERS STATE NEW JERSEY.
XX
PI Maliga P, Kuroda H, Khan MS;
XX
XX WPI: 2000-205525/18.
XX
PT New recombinant DNA constructs, for expressing high levels of
PT heterologous protein in plastids of higher plants, includes promoter, a
PT leader sequence and a downstream box element -
XX
XX Example 7; Page 69; 164pp; English.
XX
XX Primers AA61218-45 were used to construct a synthetic Streptomyces
XX hygroscopicus bar gene. The bar gene encodes phosphinothricin
XX acetyltransferase, which provides protection from phosphinothricin
XX toxicity. The synthetic gene has improved containment and enhanced
XX expression in plant plastids, and is used to produce recombinant DNA
XX constructs of the invention. The specification describes recombinant DNA
XX constructs for expressing heterologous proteins (e.g. bar gene product)
XX in the plastids of higher plants. The DNA constructs comprise a 5'
XX regulatory region which includes a promoter element, a leader sequence

CC and a downstream box element operably linked to a coding region of the
CC heterologous protein. The chimeric regulatory region enhances
CC translational efficiency of an mRNA molecule encoded by the DNA
CC construct. The DNA constructs are used for producing transformed
CC monocot and dicot plants having high levels of heterologous protein
CC expression. They can be used to drive expression of proteins with
CC agronomic, industrial or pharmaceutical importance, including production
CC of vaccines, healthcare products like human haemoglobin, industrial or
CC household enzymes. Plants which can be transformed with the constructs
CC of the invention include maize, millet, sorghum, sugar cane, rice,
CC wheat, barley, oat, rye or turf grass.
XX
SQ Sequence 40 BP; 8 A; 5 C; 9 G; 18 T; 0 other;

Query Match 14.4%; Score 13; DB 21; Length 40;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 18 aaagtctactgta 30
|||||
Db 28 aaagtctactgta 40

RESULT 24
AAV51009/C
ID AAV51009 standard; DNA; 41 BP.
XX
AC AAV51009;
XX
DT 04-JAN-1999 (first entry)
XX
DE Maize polymorphic marker S4162/G5-2 DNA.
XX
KW Polymorphic marker; allele-specific; primer; probe; amplification;
KW hybridisation; plant; hybrid certification; genetic contribution;
KW progeny; back-cross; hybrid; ancestry; maize; ss.
XX
XX Zea mays.
XX
OS
XX
FH Key Location/Qualifiers
FT 21
FT Variation /tag= a
FT /replace= "g"
FT /note= "polymorphism"
XX
PN WO9824796-A1.
XX
PD 11-JUN-1998.
XX
PF 01-DEC-1997; 97WO-US21782.
XX
PR 07-MAR-1997; 97US-0813507.
PR 02-DEC-1996; 96US-0032069.
XX
PA (AFFY-) AFFYMETRIX INC.
XX
PI Landry BS, Lemieux B, Murgineux A, Sapolsky RJ;
XX
XX WPI: 1998-333252/29.
XX
PT Brassica species allele-specific oligonucleotide probes and primers
PT - useful for plant breeding
XX
XX Claim 1; Page 43; 65pp; English.
XX
XX This DNA sequence is a region of a Zea mays genome which contains a
XX polymorphic marker. This sequence can be used in the construction of
XX allele-specific primers and probes for amplification or hybridisation,
XX e.g. to determine common or disparate ancestry between 2 or more plants,
XX to monitor the genetic contribution of an ancestral plant, to trace the
XX progeny of proprietary plants, in certification of a hybrid plant or to
XX identify the progeny of a back-crossed plant with an ancestral plant.

XX Sequence 41 BP; 10 A; 9 C; 6 G; 16 T; 0 other;
 SQ

Query Match 14.4%; Score 13; DB 19; Length 41;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatttc 17
 |||||
 DB 36 AAAAAATTATTTC 24

RESULT 25
 AAV5101/C
 ID AAV51011 standard; DNA; 41 BP.

AC AAV51011;
 XX
 DT 04-JAN-1999 (first entry)
 XX

DE Maize polymorphic marker S4IG2/G4-1 DNA.

XX Polymorphic marker; allele-specific; primer; probe; amplification;
 KW hybridisation; plant; hybrid certification; genetic contribution;
 KW progeny; back-cross; hybrid; ancestry; maize; ss.

XX Zea mays.

XX Key Location/Qualifiers
 FH variation 21
 FT /*tag= a
 FT /replace= "g"
 FT /note= "polymorphism"

XX W09924796-A1.

XX 11-JUN-1998.

XX 01-DEC-1997; 97WO-US21782.

XX 07-MAR-1997; 97US-0813507.
 PR 02-DEC-1996; 96US-0032069.

XX (AFRY-) AFFYMETRIX INC.

PI Landry BS, Lemieux B, Murigneux A, Sapolsky RJ;

DR WPI; 1998-333252/29.

XX Brassica species allele-specific oligonucleotide probes and primers
 PT - useful for plant breeding

PS Claim 1; Page 43; 65pp; English.

XX This DNA sequence is a region of a Zea mays genome which contains a
 CC polymorphic marker. This sequence can be used in the construction of
 CC allele-specific primers and probes for amplification or hybridisation,
 CC e.g. to determine common or disparate ancestry between 2 or more plants,
 CC to monitor the genetic contribution of an ancestral plant, to trace the
 CC progeny of proprietary plants, in certification of a hybrid plant or to
 CC identify the progeny of a back-crossed plant with an ancestral plant.

XX Sequence 41 BP; 12 A; 8 C; 6 G; 15 T; 0 other;
 SQ

Query Match 14.4%; Score 13; DB 19; Length 41;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatttc 17
 |||||
 DB 30 AAAAAATTATTTC 18

RESULT 26
 ID AAT25074/C
 AC AAT25074;
 XX

DT 22-OCT-1996 (first entry)

DE Human gene signature H0MGS07214.

XX Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 KW human; cloning; mapping; non-biased library; diagnosis; detection;
 KW cell typing; abnormal cell function; ss.

OS Homo sapiens.

XX W09514772-A1.

XX 01-JUN-1995.

XX 11-NOV-1994; 94WO-JP01916.

XX 12-NOV-1993; 93JP-0355504.

XX (MATS/) MATSUBARA K.
 PA (OKUBO/) OKUBO K.

PI Matsubara K, Okubo K;

DR WPI; 1995-206931/27.

XX Identifying gene signatures in 3'-directed human cDNA library - e.g.
 PT for diagnosis of abnormal cell function, by preparing cDNA that
 PT reflects relative abundance of corresp. mRNA in specific human
 PT tissues

PS Claim 1; Page 1763; 2245pp; Japanese.

XX A single-stranded DNA (or its complementary strand or the corresp.
 CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
 CC given in AAT19001-T26837 and which is able to hybridise to part of
 CC human genomic DNA, cDNA or mRNA is claimed. The GS (gene signature)
 CC sequences were obtained from 3'-directed cDNA libraries prepared
 CC from various human tissues; synthesis of cDNA was initiated from the
 CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
 CC untranslated sequence is unique to a particular mRNA species, almost
 CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
 CC is constructed so as to reflect accurately the relative abundance of
 CC different mRNAs in the particular tissue from which it was derived.
 CC The appearance frequency of a given GS in a cDNA library can be
 CC determined (esp. using primers and probes derived from the GS
 CC sequences) as a means of diagnosing abnormal cell function or for
 CC recognising different cell types.

XX Sequence 50 BP; 26 A; 3 C; 3 G; 18 T; 0 other;
 SQ

Query Match 14.4%; Score 13; DB 16; Length 50;
 Best Local Similarity 100.0%; Pred. No. 2.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 12 tatttcaagttt 24
 |||||
 DB 43 TATTCCAAGTTT 31

RESULT 27
 ID AAX52184/C
 AC AAX52184 standard; DNA; 50 BP.

XX AAX52184;

```

XX 18-JUN-1999 (first entry)
XX Synthetic plasmid synlux4 construction oligonucleotide R68.
DE DNA plasmid; lux A; lux B; Vibrio fischeri; luciferase; promoter;
XX tn9 kanamycin/neoamycin phosphotransferase; DNA synthesis;
KW replication competent double-stranded polynucleotide; ss.
XX Synthetic.
OS
XX WO9914318-A1.
XX 25-MAR-1999.
PD
XX 16-SEP-1998; 98WO-US19312.
XX 16-SEP-1997; 97US-0059017.
XX (TEXA ) UNIV TEXAS SYSTEM.
XX (TEXA ) UNIV TEXAS SYSTEM.
XX Evans GA;
XX WPI; 1999-244029/20.
XX
XX Synthesis of replication competent double-stranded polynucleotides
XX Example 4; Fig 5F; 135pp; English.
XX
XX AAX52021-212 represent oligonucleotide primers that were used to
XX construct a synthetic DNA plasmid sequence synlux4, to demonstrate the
XX method of the invention. Within the synlux4 sequence are included the
XX sequences of lux A, lux B, the A and B components of the Vibrio fischeri
XX luciferase sequence, positions of pUC19 including the origin of
XX replication and replication stability sequences, and the promoter and
XX coding sequence for tn9 kanamycin/neoamycin phosphotransferase. The
XX specification describes a method for the synthesis of replication
XX competent double-stranded polynucleotides. The method comprises
XX generating a first set of oligonucleotides corresponding to the plus
XX strand and a second set corresponding to the minus strand and
XX annealing. The method can be used for preparing polynucleotides
XX encoding sequences involved in a biochemical pathway. In particular,
XX they can be used to produce polynucleotides encoding enzymes,
XX e.g. hexokinase, phosphohexose isomerase, phosphotriokinase-1,
XX aldolase, triose-phosphate isomerase, glyceraldehyde-3-phosphate
XX dehydrogenase, phosphoglycerate kinase, phosphoglycerate mutase,
XX enolase or pyruvate kinase. They can also be used for the preparation
XX of viral particles, artificial genomes and artificial genetic systems.
XX
XX Sequence 50 BP; 19 A; 13 C; 7 G; 11 T; 0 other;
XX
XX Query Match 14.4%; Score 13; DB 20; Length 50;
XX Best Local Similarity 100.0%; Pred. No. 2.7e+03;
XX Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Oy 54 tatgttgagca 66
XX ||||||||||||
Db 44 TATGTGTGAGCA 32

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KW replication competent double-stranded polynucleotide; ss.
XX Synthetic.
OS
XX WO9914318-A1.
XX 25-MAR-1999.
PD
XX 16-SEP-1998; 98WO-US19312.
XX 16-SEP-1997; 97US-0059017.
XX (TEXA ) UNIV TEXAS SYSTEM.
XX (TEXA ) UNIV TEXAS SYSTEM.
XX Evans GA;
XX WPI; 1999-244029/20.
XX
XX Synthesis of replication competent double-stranded polynucleotides
XX Example 4; Fig 5A; 135pp; English.
XX
XX AAX52021-212 represent oligonucleotide primers that were used to
XX construct a synthetic DNA plasmid sequence synlux4, to demonstrate the
XX method of the invention. Within the synlux4 sequence are included the
XX sequences of lux A, lux B, the A and B components of the Vibrio fischeri
XX luciferase sequence, positions of pUC19 including the origin of
XX replication and replication stability sequences, and the promoter and
XX coding sequence for tn9 kanamycin/neoamycin phosphotransferase. The
XX specification describes a method for the synthesis of replication
XX competent double-stranded polynucleotides. The method comprises
XX generating a first set of oligonucleotides corresponding to the plus
XX strand and a second set corresponding to the minus strand and
XX annealing. The method can be used for preparing polynucleotides
XX encoding sequences involved in a biochemical pathway. In particular,
XX they can be used to produce polynucleotides encoding enzymes,
XX e.g. hexokinase, phosphohexose isomerase, phosphotriokinase-1,
XX aldolase, triose-phosphate isomerase, glyceraldehyde-3-phosphate
XX dehydrogenase, phosphoglycerate kinase, phosphoglycerate mutase,
XX enolase or pyruvate kinase. They can also be used for the preparation
XX of viral particles, artificial genomes and artificial genetic systems.
XX
XX Sequence 50 BP; 16 A; 7 C; 8 G; 19 T; 0 other;
XX
XX Query Match 14.4%; Score 13; DB 20; Length 50;
XX Best Local Similarity 100.0%; Pred. No. 2.7e+03;
XX Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Oy 54 tatgttgagca 66
XX ||||||||||||
Db 33 tatgttgagca 45

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RESULT 28
AAX52048
ID AAX52048 standard; DNA; 50 BP.
XX
XX AAX52048;
XX
XX 18-JUN-1999 (first entry)
XX
XX Synthetic plasmid synlux4 construction oligonucleotide F28.
XX
XX DNA plasmid; lux A; lux B; Vibrio fischeri; luciferase; promoter;
KW tn9 kanamycin/neoamycin phosphotransferase; DNA synthesis;
KW

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RESULT 29
AAX70046/C
ID AAX70046 standard; RNA; 17 BP.
XX
XX AAX70046;
XX
XX 28-JUL-1999 (first entry)
XX
XX Human flt1 VEGF receptor hammerhead ribozyme substrate #1341.
XX
XX Vascular endothelial growth factor receptor; flt-1;
KW flk-1; KDR; hammerhead ribozyme; hairpin ribozyme; cleavage;
KW tumour angiogenesis; psoriasis; rheumatoid arthritis; ocular disease;
KW fms-like tyrosine kinase 1; kinase insert domain containing receptor;
KW foetal liver kinase 1; ss.
XX
XX Homo sapiens.
OS
XX
XX W09715662-A2.
PN

```

XX 01-MAY-1997.
 PD 25-OCT-1996; 96WO-US17480.
 XX
 XX 11-JAN-1996; 96US-0584040.
 PR 26-OCT-1995; 95US-0005974.
 XX
 PA (CHIR) CHIRON CORP.
 PA (RIBO-) RIBOZYME PHARM INC.
 XX
 PI Escobedo J, McSwigen J, Pavco P, Stinchcomb D;
 DR WPI; 1997-259017/23.
 XX
 PT Nucleic acid molecule modulating VEGF receptor(s) gene expression or
 PT mRNA stability - useful for treating e.g. tumour angiogenesis,
 PT psoriasis, rheumatoid arthritis, etc., in a human patient
 XX
 PS Claim 4; Page 87; 218pp; English.
 XX
 CC The present invention describes nucleic acid molecules which modulate
 CC the synthesis, expression and/or stability of a mRNA encoding 1 or more
 CC receptors of vascular endothelial growth factor (VEGF). A patient
 CC (preferably human) having a condition associated with the level of the
 CC fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing
 CC receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour
 CC angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can
 CC be treated by administering the nucleic acid molecule or the expression
 CC vector to the patient. AAX67275 to AAX75752 represent specific examples
 CC of nucleic acid molecules from the present invention.
 XX
 SQ Sequence 17 BP; 8 A; 1 C; 4 G; 4 U; 0 other;

Query Match 13.3%; Score 12; DB 18; Length 17;
 Best Local Similarity 100.0%; Pred. NO. 8.3e+03;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 13 attcaaatgtt 24
 |||||
 DB 15 ATTTCAAAGTTT 4

RESULT 30
 AAX70047/C
 ID AAX70047 standard; RNA; 17 BP.
 XX
 AC AAX70047;
 XX
 DT 28-JUL-1999 (first entry)
 XX
 DE Human flt1 VEGF receptor hammerhead ribozyme substrate #1342.
 XX
 KW Vascular endothelial growth factor receptor; VEGF receptor; flt-1;
 KW flk-1; KDR; hammerhead ribozyme; hairpin ribozyme; cleavage;
 KW tumour angiogenesis; psoriasis; rheumatoid arthritis; ocular disease;
 KW fms-like tyrosine kinase 1; kinase insert domain containing receptor;
 KW foetal liver kinase 1; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO9715662-A2.
 XX
 PD 01-MAY-1997.
 XX
 PF 25-OCT-1996; 96WO-US17480.
 XX
 PR 11-JAN-1996; 96US-0584040.
 PR 26-OCT-1995; 95US-0005974.
 XX
 PA (CHIR) CHIRON CORP.
 PA (RIBO-) RIBOZYME PHARM INC.

XX Escobedo J, McSwigen J, Pavco P, Stinchcomb D;
 XX WPI; 1997-259017/23.
 DR
 XX
 PT Nucleic acid molecule modulating VEGF receptor(s) gene expression or
 PT mRNA stability - useful for treating e.g. tumour angiogenesis,
 PT psoriasis, rheumatoid arthritis, etc., in a human patient
 XX
 PS Claim 4; Page 87; 218pp; English.

XX The present invention describes nucleic acid molecules which modulate
 CC the synthesis, expression and/or stability of a mRNA encoding 1 or more
 CC receptors of vascular endothelial growth factor (VEGF). A patient
 CC (preferably human) having a condition associated with the level of the
 CC fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing
 CC receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour
 CC angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can
 CC be treated by administering the nucleic acid molecule or the expression
 CC vector to the patient. AAX67275 to AAX75752 represent specific examples
 CC of nucleic acid molecules from the present invention.

SQ Sequence 17 BP; 8 A; 1 C; 3 G; 5 U; 0 other;

Query Match 13.3%; Score 12; DB 18; Length 17;
 Best Local Similarity 100.0%; Pred. NO. 8.3e+03;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 13 attcaaatgtt 24
 |||||
 DB 14 ATTTCAAAGTTT 3

RESULT 31
 AAA21453
 ID AAA21453 standard; RNA; 17 BP.

AC AAA21453;

DT 19-JUN-2000 (first entry)

DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4679.

XX Human; aryl hydrocarbon nuclear transport; ARNT; TIE-2; angiogenesis;
 KW integrin alpha 6 subunit; integrin subunit beta 3; hairpin ribozyme;
 KW hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
 KW ophthalmological; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
 KW dermatological; RNA cleavage; cancer; diabetic retinopathy; arthritis;
 KW age related macular degeneration; inflammation; neovascular glaucoma;
 KW myopic degeneration; psoriasis; verruca vulgaris; angiofibroma;
 KW tuberosus sclerosis; pot-wine stain; Sturge Weber syndrome;
 KW Kippel-Trenaunay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.

XX Homo sapiens.

PN WO9950403-A2.

PD 07-OCT-1999.

PF 24-MAR-1999; 99WO-US06507.

PR 27-MAR-1998; 98US-0079678.

PA (RIBO-) RIBOZYME PHARM INC.

XX Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwigen JA;

DR WPI; 1999-591315/50.

PT Novel ribozymes for modulating the synthesis, expression and/or
 stability of an mRNA encoding an angiogenic factors

PS Claim 55; Page 209; 305pp; English.

XX The present invention describes enzymatic cleave RNA encoded by an aryl
 CC RNA cleaving activity, which specifically cleave RNA encoded by an aryl
 CC hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
 CC gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
 CC AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
 CC and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
 CC corresponding target sequences. AAA17685 to AAA18385 and AAA19087 to
 CC AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
 CC and AAA19155 to AAA19222 represent their corresponding target sequences;
 CC AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
 CC sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
 CC AAA21596 to AAA21688 represent their corresponding target sequences;
 CC AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequence
 CC for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
 CC AAA23422 represent their corresponding target sequences. The ribozymes of
 CC the invention are used for modulating the synthesis, expression and/or
 CC stability of an mRNA encoding angiogenic factor, especially ARNT,
 CC integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
 CC especially used to treat cancer, diabetic retinopathy, age related
 CC macular degeneration (ARMD), inflammation, and arthritis, as well as
 CC neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
 CC angioidioma of tuberous sclerosis, pot-wine stains, Sturge Weber
 CC syndrome, Kippel-Trennauay-Weber syndrome, Osler-Weber-Rendu syndrome,
 CC and other syndromes and diseases related to the levels of ARNT, Tie-2,
 CC integrin subunit alpha-6, or integrin subunit beta-3.
 CC
 XX Sequence 17 BP; 6 A; 1 C; 2 G; 8 U; 0 other;

Query Match 13.3%; Score 12; DB 20; Length 17;
 Best Local Similarity 58.3%; Pred. No. 8.3e+03;
 Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Oy 7 aaatatttca 18
 |||||:::||||
 Db 4 aaauuuuuca 15

RESULT 32
 AAA21454
 ID AAA21454 standard; RNA; 17 BP.
 XX
 AC AAA21454;

XX 19-JUN-2000 (first entry)

DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4680.

XX Human: aryl hydrocarbon nuclear transport; ARNT; Tie-2; angiogenesis;
 KW Integrin alpha 6 subunit; Integrin subunit beta 3; hairpin ribozyme;
 KW hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
 KW ophthalmologic; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
 KW dermatological; RNA cleavage; cancer; diabetic retinopathy; arthritis;
 KW age related macular degeneration; inflammation; neovascular glaucoma;
 KW myopic degeneration; psoriasis; verruca vulgaris; angioidioma;
 KW tuberous sclerosis; pot-wine stain; Sturge Weber syndrome;
 KW Kippel-Trennauay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.

XX Homo sapiens.

OS W09950403-A2.

PN 07-OCT-1999.

XX 24-MAR-1999; 99WO-US06507.

XX 27-MAR-1998; 98US-0079678.

XX (RIBO-) RIBOZYME PHARM INC.

PA Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwiggen JA;

XX WPI; 1999-591315/50.

DR Novel ribozymes for modulating the synthesis, expression and/or
 XX stability of an mRNA encoding an angiogenic factors
 PT
 XX
 PS Claim 55; Page 209; 305pp; English.

CC The present invention describes enzymatic cleave RNA encoded by an aryl
 CC RNA cleaving activity, which specifically cleave RNA encoded by an aryl
 CC hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
 CC gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
 CC AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
 CC and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
 CC corresponding target sequences. AAA17685 to AAA18385 and AAA19087 to
 CC AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
 CC and AAA19155 to AAA19222 represent their corresponding target sequences;
 CC AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
 CC sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
 CC AAA21596 to AAA21688 represent their corresponding target sequences;
 CC AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequence
 CC for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
 CC AAA23422 represent their corresponding target sequences. The ribozymes of
 CC the invention are used for modulating the synthesis, expression and/or
 CC stability of an mRNA encoding angiogenic factor, especially ARNT,
 CC integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
 CC especially used to treat cancer, diabetic retinopathy, age related
 CC macular degeneration (ARMD), inflammation, and arthritis, as well as
 CC neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
 CC angioidioma of tuberous sclerosis, pot-wine stains, Sturge Weber
 CC syndrome, Kippel-Trennauay-Weber syndrome, Osler-Weber-Rendu syndrome,
 CC and other syndromes and diseases related to the levels of ARNT, Tie-2,
 CC integrin subunit alpha-6, or integrin subunit beta-3.
 CC
 XX Sequence 17 BP; 7 A; 1 C; 2 G; 7 U; 0 other;

Query Match 13.3%; Score 12; DB 20; Length 17;
 Best Local Similarity 58.3%; Pred. No. 8.3e+03;
 Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Oy 7 aaatatttca 18
 |||||:::||||
 Db 3 aaauuuuuca 14

RESULT 33
 AAA21455
 ID AAA21455 standard; RNA; 17 BP.
 XX
 AC AAA21455;

XX 19-JUN-2000 (first entry)

DE Integrin alpha 6 subunit substrate sequence SEQ ID NO:4681.

XX Human: aryl hydrocarbon nuclear transport; ARNT; Tie-2; angiogenesis;
 KW Integrin alpha 6 subunit; Integrin subunit beta 3; hairpin ribozyme;
 KW hammerhead ribozyme; angiogenic factor; cytoskeletal; antidiabetic;
 KW ophthalmologic; antiinflammatory; antiarthritic; antipsoriatic; ARMD;
 KW dermatological; RNA cleavage; cancer; diabetic retinopathy; arthritis;
 KW age related macular degeneration; inflammation; neovascular glaucoma;
 KW myopic degeneration; psoriasis; verruca vulgaris; angioidioma;
 KW tuberous sclerosis; pot-wine stain; Sturge Weber syndrome;
 KW Kippel-Trennauay-Weber syndrome; Osler-Weber-Rendu syndrome; ss.

XX Homo sapiens.

OS W09950403-A2.

PN 07-OCT-1999.

XX 24-MAR-1999; 99WO-US06507.

XX 27-MAR-1998: 98US-0079678.
 PR (RIBO-) RIBOZYME PHARM INC.
 XX PA
 XX PI Pavco PA, Roberts E, Jarvis T, Coesholt C, McSwiggen JA;
 XX WPI: 1999-591315/50.
 DR
 XX Novel ribozymes for modulating the synthesis, expression and/or
 PT stability of an mRNA encoding an angiogenic factors -
 PT
 XX
 PS Claim 55; Page 209; 305pp; English.
 XX
 CC The present invention describes enzymatic nucleic acid molecules with
 CC RNA cleaving activity, which specifically cleave RNA encoded by an aryl
 CC hydrocarbon nuclear transporter (ARNT) gene, an integrin subunit beta 3
 CC gene, an integrin alpha 6 subunit gene, or a Tie-2 gene. AAA16775 to
 CC AAA17167 and AAA17561 to AAA17622 represent ribozyme sequences for ARNT,
 CC and AAA17168 to AAA17560 and AAA17623 to AAA17684 represent their
 CC corresponding target sequences; AAA17685 to AAA18385 and AAA19087 to
 CC AAA19154 represent ribozyme sequences for Tie-2, and AAA18386 to AAA19086
 CC and AAA19155 to AAA19222 represent their corresponding target sequences;
 CC AAA19223 to AAA20361 and AAA21501 to AAA21595 represent ribozyme
 CC sequences for integrin alpha 6 subunit, and AAA20362 to AAA21500 and
 CC AAA21596 to AAA21688 represent their corresponding target sequences;
 CC AAA21689 to AAA22475 and AAA23263 to AAA23342 represent ribozyme sequences
 CC for integrin subunit beta 3, and AAA22476 to AAA23262, AAA23343 to
 CC AAA23422 represent their corresponding target sequences. The ribozymes of
 CC the invention are used for modulating the synthesis, expression and/or
 CC stability of an mRNA encoding angiogenic factor, especially ARNT,
 CC integrin subunit beta-3, integrin subunit alpha-6, or Tie-2. They are
 CC especially used to treat cancer, diabetic retinopathy, age related
 CC macular degeneration (AMD), inflammation, and arthritis, as well as
 CC neovascular glaucoma, myopic degeneration, psoriasis, verruca vulgaris,
 CC syngiofibroma of tuberous sclerosis, pot-wine stains, Sturge Weber
 CC syndrome, Kippel-Trenaunay-Weber syndrome, Osler-Weber-Rendu syndrome,
 CC and other syndromes and diseases related to the levels of ARNT, Tie-2,
 CC integrin subunit alpha-6, or integrin subunit beta-3.
 CC
 XX Sequence 17 BP; 7 A; 1 C; 1 G; 8 U; 0 other;
 SQ

Query Match 13.3%; Score 12; DB 20; Length 17;
 Best Local Similarity 58.3%; Pred. No. 8.3e+03;
 Matches 7; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY 7 aaaaattattca 18
 |||||:|||||
 Db 1 aaauuuuuuca 12

RESULT 34
 AAF03088/C
 ID AAF03088 standard; DNA; 17 BP.
 XX
 AC AAF03088;
 XX
 DT 16-FEB-2001 (first entry)
 XX
 DE Hammerhead ribozyme substrate #1383.
 XX
 KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
 KM interferon alpha; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200061729-A2.
 XX
 PD 19-OCT-2000.
 XX
 PF 11-APR-2000; 2000WO-US09721.
 XX
 PS

PR 12-APR-1999: 99US-0129390.
 XX
 XX (RIBO-) RIBOZYME PHARM INC.
 XX PA
 XX PI Blatt L, Zwick M, Pavco P, McSwiggen J;
 XX WPI: 2000-647423/62.
 DR
 XX Enzymatic and antisense nucleic acid inhibition of repressor genes,
 PT useful for producing e.g. granulocyte colony stimulating factor
 PT protein, interferon alpha and erythropoietin -
 PT
 XX
 PS Claim 37; Page 87; 164pp; English.
 XX
 CC The present invention relates to enzymatic and antisense nucleic acid
 CC molecules that act as inhibitors of the expression of repressor genes
 CC encoding the TR2 Orphan receptor, EAR3/CODP-TF-1, the GATA
 CC transcription factor gene, IRF-2 and/or the CAA17 Displacement
 CC protein (CDP). Inhibition of the repressors removes prevents
 CC inhibition (and consequently increases expression of) genes involved in
 CC the production of erythropoietin, granulocyte colony stimulating factor
 CC protein and interferon alpha.
 CC
 XX Sequence 17 BP; 6 A; 1 C; 0 G; 10 T; 0 other;
 SQ

Query Match 13.3%; Score 12; DB 21; Length 17;
 Best Local Similarity 100.0%; Pred. No. 8.3e+03;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
 |||||:|||||
 Db 17 AAAAAATTATT 6

RESULT 35
 AAF03089/C
 ID AAF03089 standard; DNA; 17 BP.
 XX
 AC AAF03089;
 XX
 DT 16-FEB-2001 (first entry)
 XX
 DE Hammerhead ribozyme substrate #1384.
 XX
 KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
 KM interferon alpha; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200061729-A2.
 XX
 PD 19-OCT-2000.
 XX
 PF 11-APR-2000; 2000WO-US09721.
 XX
 PR 12-APR-1999; 99US-0129390.
 XX
 PA (RIBO-) RIBOZYME PHARM INC.
 XX
 PI Blatt L, Zwick M, Pavco P, McSwiggen J;
 XX
 XX WPI: 2000-647423/62.
 DR
 XX Enzymatic and antisense nucleic acid inhibition of repressor genes,
 PT useful for producing e.g. granulocyte colony stimulating factor
 PT protein, interferon alpha and erythropoietin -
 PT
 XX
 PS Claim 37; Page 87; 164pp; English.
 XX
 CC The present invention relates to enzymatic and antisense nucleic acid
 CC molecules that act as inhibitors of the expression of repressor genes
 CC encoding the TR2 Orphan receptor, EAR3/CODP-TF-1, the GATA

transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 8 A; 1 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 14 AAAAAATTATT 3

RESULT 36
AA03090/c
ID AAF03090 standard; DNA: 17 BP.

AC AAF03090;
DT 16-FEB-2001 (first entry)

DE Hammerhead ribozyme substrate #1385.

KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
KM Interferon alpha; ss.

OS Homo sapiens.

XX WO200061729-A2.

XX 19-OCT-2000.

XX 11-APR-2000; 2000WC-US09721.

XX 12-APR-1999; 99US-0129390.

PA (RIBO-) RIBOZYME PHARM INC.

PI Blatt L, Zwick M, Pavco P, MCSwigen J;

WPI: 2000-647423/62.

Enzymatic and antisense nucleic acid inhibition of repressor genes,
useful for producing e.g. granulocyte colony stimulating factor
protein, interferon alpha and erythropoietin -

Claim 37; Page 87; 164pp; English.

The present invention relates to enzymatic and antisense nucleic acid molecules that act as inhibitors of the expression of repressor genes encoding the IR2 Orphan receptor, EAR3/COUP-1, the GATA transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 8 A; 1 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 13 AAAAAATTATT 2

RESULT 37
AA03091/c
ID AAF03091 standard; DNA: 17 BP.

AC AAF03091;

DT 16-FEB-2001 (first entry)

DE Hammerhead ribozyme substrate #1386.

KW Ribozyme; erythropoietin; granulocyte colony stimulating factor;
KM Interferon alpha; ss.

OS Homo sapiens.

XX WO200061729-A2.

XX 19-OCT-2000.

XX 11-APR-2000; 2000WC-US09721.

XX 12-APR-1999; 99US-0129390.

PA (RIBO-) RIBOZYME PHARM INC.

PI Blatt L, Zwick M, Pavco P, MCSwigen J;

WPI: 2000-647423/62.

Enzymatic and antisense nucleic acid inhibition of repressor genes,
useful for producing e.g. granulocyte colony stimulating factor
protein, interferon alpha and erythropoietin -

Claim 37; Page 87; 164pp; English.

The present invention relates to enzymatic and antisense nucleic acid molecules that act as inhibitors of the expression of repressor genes encoding the IR2 Orphan receptor, EAR3/COUP-1, the GATA transcription factor gene, IRF-2 and/or the CATT Displacement Protein (CDP). Inhibition of the repressors removes prevents inhibition (and consequently increases expression of) genes involved in the production of erythropoietin, granulocyte colony stimulating factor protein and interferon alpha.

Sequence 17 BP; 9 A; 0 C; 0 G; 8 T; 0 other;

Query Match 13.3%; Score 12; DB 21; Length 17;
Best Local Similarity 100.0%; Pred. No. 8.3e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaattatt 15
|||||
DB 12 AAAAAATTATT 1

RESULT 38
AAA08921
ID AAA08921 standard; DNA: 18 BP.

AC AAA08921;

DT 01-AUG-2000 (first entry)

DE Human survivin DNA antisense oligonucleotide, ISIS 23663.

KW Survivin; inhibitor of apoptosis; IAP; caspase inhibitor; caspase-3;
cell cycle regulation; cancer; cytostatic; antisense oligonucleotide; ss.

OS Synthetic.
XX Homo sapiens.

Sequence 20 BP; 8 A; 4 C; 3 G; 5 T; 0 other;

Query Match	13.3%	Score 12;	DB 11;	Length 20;
Best Local Similarity	100.0%	Pred. NO. 8.3e+03;		
Matches 12;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

Oy	74	caatgaaagcaa	85
Db	2	caatgaaagcaa	13

```

RESULT 43
AAT80813/c
ID AAT80813 standard; cDNA; 20 BP

```

AC AAT80813;

DT 14-APR-1998 (first entry)

Staphylococcus aureus Gene #22 PCR primer sequence 4.

KW Staphylococcus aureus WCUH 29; antagonist; antibacterial;
KM vaccine; disease; protection; isolation; PCR primer; ss.

05 Synthetic.

05 Staphylococcus aureus

PN WO9731114-A2.

PD 28-AUG-1997.

PF 25-FEB-1997; 97WO-GB00524

PR 26-FEB-1996; 96GB-0004045

PA (SMIK) SMITHKLINE BEECHAM PLC.

PI Burnham MKR, Hodgson JE,

DR WPI; 1997-435166/40

PT New *Staphylococcus aureus* polynucleotide and polypeptide(s) - for
isolating antagonist of the polypeptide(s) useful as anti-bacterials
XX
PS Disclosure, Page 52, 117pp, English.

The present sequence represents a PCR primer used in the present invention describing novel polypeptides, which can optionally be expressed in NCIMB 40771. The polypeptides, and polynucleotides encoding it, are derived from *Staphylococcus aureus*, cells expressing ligands binding the polypeptides can be used to isolated candidate compounds that bind and inhibit the activity of the polypeptides. Such compounds can be used as anti-bacterial compounds. The polypeptides may also be used as immunogens to vaccinate an animal for protection against *Staphylococcus aureus* caused disease.

Sequence 20 BP; 11 A; 4 C; 2 G; 3 T; 0 other;

Query Match	13.3%;	Score 12;	DB 18;	Length 20;
Best Local Similarity	100.0%;	Pred. NO. 8.3e+03;		
Matches 12;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

```

0y      39 gtttcacgttt 50
          |||||
Db      20 GTTTTCATGTTT 9

```

RESULT	44
AAV72697/c	
ID	AAV72697 standard; DNA; 20 BP

AC	AAV72697;
XX	
DT	17-FEB-1999 (first entry)

Corn kernel oil concentration controlling loci marker s1384 primer 2.

KW Corn; kernel oil; concentration; trait controlling loci; genetic marker;
ZW Zee maw; brooding; BCD primer; ss

KW Zea mays; breeding; PCR primer; ss.

OS Synthetic

05 Zea mays.

PN WO9842870-A1.

PD 01-OCT-1998

PF 19-MAR-1998; 98WO-US05550

PR 24-MAR-1997; 97US-0041515

PA (DUPO) DU PONT DE NEMOURS & CO E. I.

PI Reiter RS

DR WPI; 1998-609896/51.

PT Breeding corn with increased oil concentration - comprises use of
PT genetic markers to identify trait loci controlling kernel oil
PT concentration
XX
XS Example 2; Page 5; 50pp; English.

A new method has been developed of breeding to corn with increased kernel oil concentration. The method comprises: (a) selecting a corn plant from a breeding population using at least one of the genetic markers s1375, s1384, s1394, s1416, s1422, s1432, s1457, s1480, s1476, s1478, s1484, s1500, s1513, s1539, s1544, s1545, s1630, s1633, s1647, s1750, s1756, s1757, s1767, s1772, s1774, s1780, s1797, s1813, s1816, s1817, s1836, s1853, s1860, s1870, s1921, s1922, s1925, s1931, s1933, s1939, s1946, s1949, s2054, s2055, s2057, s2058, s2097, s2122, s2125, s2150, s2156, and s2175; and (b) crossing the selected plant with a second plant and obtaining progeny with increased kernel oil concentration. Also described are: (1) a method for identifying corn plants or lines for use as parents to create a breeding population, comprising: (a) genotyping corn plants or lines with one or more of the above genetic markers; and (b) identifying plants or lines which are predicted to produce transgressive segregants for kernel oil concentration; and (2) trait loci controlling kernel oil concentration mapped by the above genetic markers, with the exception of s1480. AAT2694 to AAT2757 represent PCR primers which are used to amplify the genetic markers for use in the method of the invention.

CC AAV72694 to AAV72797 represent PCR primers which are used to amplify the
CC genetic markers for use in the method of the invention.

Sequence 20 BP; 4 A; 7 C; 3 G; 6 T; 0 other,

Query Match	13.38;	Score 12;	DB 19;	Length 20;
Best Local Similarity	100.0%;	Pred. No. 8.3e+03;		
Matches 12;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

QY	77	tcgaagcaagt	88
Db	12	TGAAGCAAGTC	1

RESULT	45
AAX92327	
ID	AAX92327 standard; DNA; 20 BP.

AC AAX92327;

DT 13-SEP-1999 (first entry)

DE PCR primer used to amplify an ORF of *Chlamydia pneumoniae*

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
 KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
 KW vaccine; neutralising epitope; PCR primer; ss.

OS Synthetic.
 OS Chlamydia pneumoniae.

PM W09927105-A2.

PD 03-JUN-1999.

PF 20-NOV-1998; 98WO-1B01890.

PR 04-NOV-1998; 98US-0107078.

PR 21-NOV-1997; 97ER-0014673.

PA (GSET) GENSET.

PI Griffiths R;

DR WPI; 1999-357842/30.

Genome sequence of Chlamydia pneumoniae

Page 1503; Disclosure; 1912pp; English.

CC AAY91991-X97517 represent PCR primers used to amplify open reading
 CC frames and other nucleic acid sequences from the genome of
 CC Chlamydia pneumoniae (see AAY91990). C. pneumoniae causes respiratory
 CC disease such as pneumonia and bronchitis and is thought to be a
 CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent
 CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded
 CC by the open reading frames of the C. pneumoniae genome (see AAY34584-
 CC AAY35879) can be used in immunogenic compositions as vaccines. Vectors
 CC containing C. pneumoniae nucleotides sequences can also be used as
 CC immunogenic compositions, especially where the vector directs the
 CC expression of a neutralising epitope of C. pneumoniae.

SO Sequence 20 BP; 3 A; 5 C; 5 G; 7 T; 0 other;

Query Match 13.3%; Score 12; DB 20; Length 20;

Best Local Similarity 100.0%; Pred. No. 8.3e+03;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 atgttttcattgt 48

Db 9 atgttttcattgt 20

Search completed: January 24, 2002, 03:28:21
 Job time: 3673 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:22:33 : Search time 93.51 Seconds
(without alignments)
791.983 Million cell updates/sec

Title: US-09-531-438-3

Perfect score: 327

Sequence: 1 atttggaatcttaattt.....ttcatgtttctattgtt 327

Scoring table: OLIGO_MUC

Gapop 60.0 , Gapext 60.0

Searched: 351203 seqs, 113238999 residues

Word size : 0

Total number of hits satisfying chosen parameters: 495388

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database :

Issued_Patents_NA:*
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2: /cgn2_6/prodata/2/ina/5B_COMB.seq:*
3: /cgn2_6/prodata/2/ina/6A_COMB.seq:*
4: /cgn2_6/prodata/2/ina/6B_COMB.seq:*
5: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:*
6: /cgn2_6/prodata/2/ina/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	4.9	29	3	US-08-816-977-12 Sequence 12, Appl
2	16	4.9	36	1	US-08-629-600-16 Sequence 16, Appl
3	15	4.6	27	1	US-08-120-827-66 Sequence 66, Appl
4	15	4.6	27	1	US-08-478-675-66 Sequence 66, Appl
5	15	4.6	30	5	PCT-US92-10792-3 Sequence 3, Appl1
6	15	4.6	32	1	US-08-256-261-29 Sequence 29, Appl
7	15	4.6	32	3	US-08-852-299-29 Sequence 29, Appl
8	14	4.3	18	2	US-09-205-204-20 Sequence 20, Appl
9	14	4.3	29	3	US-08-816-977-12 Sequence 12, Appl
10	14	4.3	30	2	US-08-629-001A-31 Sequence 31, Appl
11	14	4.3	30	4	US-08-642-274D-110 Sequence 110, App
12	14	4.3	36	1	US-08-629-600-16 Sequence 16, Appl
13	14	4.3	37	2	US-08-403-853-8 Sequence 8, Appl1
14	13	4.0	18	3	US-08-847-844A-113 Sequence 113, App
15	13	4.0	18	4	US-08-686-968C-113 Sequence 13, Appl
16	13	4.0	20	3	US-09-288-461-79 Sequence 79, Appl
17	13	4.0	21	3	US-08-691-045-61 Sequence 61, Appl
18	13	4.0	24	3	US-08-672-215-1 Sequence 1, Appl1
19	13	4.0	28	1	US-08-120-827-64 Sequence 64, Appl
20	13	4.0	28	1	US-08-478-675-64 Sequence 64, Appl
21	13	4.0	30	2	US-08-629-001A-79 Sequence 79, Appl
22	13	4.0	30	4	US-08-642-274D-158 Sequence 158, App
23	13	4.0	31	1	US-08-330-638D-5 Sequence 5, Appl1
24	13	4.0	31	2	US-08-906-746A-5 Sequence 5, Appl1
25	13	4.0	36	1	US-08-247-809A-14 Sequence 14, Appl
26	13	4.0	36	2	US-08-711-728-14 Sequence 14, Appl
27	13	4.0	37	2	US-08-097-554A-45 Sequence 45, Appl

28	13	4.0	37	2	US-08-484-575A-10 Sequence 10, Appl
29	13	4.0	37	3	US-08-477-459-10 Sequence 10, Appl
30	13	4.0	37	3	US-08-480-640A-45 Sequence 45, Appl
31	13	4.0	37	3	US-08-479-869-10 Sequence 10, Appl
32	13	4.0	37	3	US-08-295-802-45 Sequence 45, Appl
33	13	4.0	37	3	US-08-486-414-10 Sequence 10, Appl
34	13	4.0	37	3	US-08-488-237A-45 Sequence 45, Appl
35	13	4.0	37	5	PCT-US94-01826A-10 Sequence 10, Appl
36	13	4.0	37	5	PCT-US94-02252A-10 Sequence 10, Appl
37	13	4.0	38	1	PCT-US96-00547-40 Sequence 40, Appl
38	13	4.0	39	1	US-08-105-483-168 Sequence 168, App
39	13	4.0	39	1	US-08-709-209-168 Sequence 56, Appl
40	13	4.0	39	1	US-08-303-275-56 Sequence 168, App
41	13	4.0	39	2	US-08-458-101-168 Sequence 168, App
42	13	4.0	39	2	US-09-028-361A-19 Sequence 19, Appl
43	13	4.0	40	1	US-08-199-507B-39 Sequence 39, Appl
44	13	4.0	40	1	US-08-441-828-39 Sequence 39, Appl
45	13	4.0	41	3	US-08-930-503A-11 Sequence 11, Appl

ALIGNMENTS

RESULT 1
US-08-816-977-12
; Sequence 12, Application US/08816977
; Patent No. 6080400
; GENERAL INFORMATION:
; APPLICANT: Williams, James A.
; APPLICANT: Byrne, Lisa M.
; APPLICANT: Pugh, Charles S.G.
; TITLE OF INVENTION: Prevention And Treatment Of
; TITLE OF INVENTION: Verotoxin-Induced Disease
; NUMBER OF SEQUENCES: 49
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Medlen & Carroll, LLP
; STREET: 220 Montgomery Street, Suite 2200
; CITY: San Francisco
; STATE: California
; COUNTRY: United States of America
; ZIP: 94104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US-08-816-977
; FILING DATE: 13-MAR-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MacKnight, Kamrin T.
; REGISTRATION NUMBER: 38,230
; REFERENCE/DOCKET NUMBER: OPMD-02450
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 705-8410
; TELEFAX: (415) 397-8338
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-816-977-12

Query Match 4.9%; Score 16; DB 3; Length 29;
Best Local Similarity 100.0%; Pred. No. 3; Ie+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 189 aaataatttttta 204
|||||

Db 9 AAAATATTATTTTTA 24

RESULT 2

US-08-629-600-16

Sequence 16, Application US/08629600

Patent No. 5783196

GENERAL INFORMATION:

APPLICANT: NORIEGA, Fernando

APPLICANT: LEVINE, Myron M.

TITLE OF INVENTION: GDA MUTANTS OF SHIGELLA

TITLE OF INVENTION: AND VACCINES CONTAINING THE SAME

NUMBER OF SEQUENCES: 18

CORRESPONDENCE ADDRESS:

ADDRESS: SUGHRUE, MION, ZINN, MACPEAK & SEAS

STREET: 2100 Pennsylvania Avenue, N.W., Suite 800

CITY: Washington, D.C.

STATE: D.C.

COUNTRY: U.S.A.

ZIP: 20037

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/629,600

FILING DATE: 9-APR-1996

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: KIT, Gordon

REGISTRATION NUMBER: 30,764

REFERENCE/DOCKET NUMBER: A-6765

TELEPHONE: (202) 293-7060

TELEFAX: (202) 293-7860

INFORMATION FOR SEQ ID NO: 16:

SEQUENCE CHARACTERISTICS:

LENGTH: 36 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA

HYPOTHETICAL: NO

US-08-629-600-16

Query Match 4.9%; Score 16; DB 1; Length 36;

Best Local Similarity 100.0%; Pred. No. 3e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204

Db 14 AAAATATTATTTTTA 29

RESULT 3

US-08-120-827-66

Sequence 66, Application US/08120827

Patent No. 5525495

GENERAL INFORMATION:

APPLICANT: KEENE, JACK D.

APPLICANT: KING, PETER H.

APPLICANT: LEVINE, TODD

TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE

TITLE OF INVENTION: RECOGNITION, BINDING AND EXPRESSION OF THE RIBONUCLEIC ACIDS

NUMBER OF SEQUENCES: 101

CORRESPONDENCE ADDRESS:

ADDRESS: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESS: P.C.

STREET: 1755 Jefferson Davis Highway, Fourth Floor

CITY: Arlington

STATE: Virginia

COUNTRY: U.S.A.

ZIP: 22202

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/120,827

FILING DATE: 15-SEP-1993

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Oblon, No. 5525495man F.

REGISTRATION NUMBER: 24,618

REFERENCE/DOCKET NUMBER: 714-158-0 CIP

TELEPHONE: (703)413-3000

TELEFAX: (703)413-2220

INFORMATION FOR SEQ ID NO: 66:

SEQUENCE CHARACTERISTICS:

LENGTH: 27 base pairs

TYPE: nucleic acid

STRANDEDNESS: unknown

TOPOLOGY: unknown

MOLECULE TYPE: RNA (genomic)

US-08-120-827-66

Query Match

Best Local Similarity 4.6%; Score 15; DB 1; Length 27;

Matches 5; Conservative 10; Mismatches 0; Indels 0; Gaps 0;

OY 197 tattttattttaaa 211

Db 12 UAUUUUUUUUUUAAA 26

RESULT 4

US-08-478-675-66

Sequence 66, Application US/08478675

Patent No. 5773246

GENERAL INFORMATION:

APPLICANT: KEENE, JACK D.

APPLICANT: KING, PETER H.

TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE

TITLE OF INVENTION: RECOGNITION, BINDING AND EXPRESSION OF THE RIBONUCLEIC ACIDS

NUMBER OF SEQUENCES: 101

CORRESPONDENCE ADDRESS:

ADDRESS: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESS: P.C.

STREET: 1755 Jefferson Davis Highway, Fourth Floor

CITY: Arlington

STATE: Virginia

COUNTRY: U.S.A.

ZIP: 22202

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/478,675

FILING DATE: 07-JUN-1996

CLASSIFICATION: 536

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/120,827

FILING DATE: 15-SEP-1993

ATTORNEY/AGENT INFORMATION:

NAME: Oblon, No. 5773246man F.

REGISTRATION NUMBER: 24,618
REFERENCE/DOCKET NUMBER: 714-158-0 CIP
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)413-3000
TELEFAX: (703)413-2220
TELEX: 248855 OPAT UR
INFORMATION FOR SEQ ID NO: 66:
SEQUENCE CHARACTERISTICS:
LENGTH: 27 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: RNA (genomic)
US-08-478-675-66

Query Match 4.6%; Score 15; DB 1; Length 27;
Best Local Similarity 33.3%; Pred. No. 7.9e+02;
Matches 5; Conservative 10; Mismatches 0; Indels 0; Gaps 0;

QY 197 tattttattttaa 211
:|:::|:::|:::|:::|
Db 12 UAUUUUAUUUUAAA 26

RESULT 5
PCT-US92-10792-3
Sequence 3, Application PC/TUS9210792
GENERAL INFORMATION:
APPLICANT: Jayasena, Sumedha D.
APPLICANT: Johnston, Brian H.
TITLE OF INVENTION: Triple Helix Formation at
NUMBER OF SEQUENCES: 53
CORRESPONDENCE ADDRESS:
ADDRESSEE: SRI International
STREET: 333 Ravenswood Avenue
CITY: Menlo Park
STATE: CA
COUNTRY: USA
ZIP: 94025
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/10792
FILING DATE: 19921211
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/826,934
FILING DATE: 21-JAN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/808,452
FILING DATE: 13-DEC-1991
ATTORNEY/AGENT INFORMATION:
NAME: Fabian, Gary R.
REGISTRATION NUMBER: 33,875
REFERENCE/DOCKET NUMBER: P-3141
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 859-4550
TELEFAX: (415) 859-3880
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: OLIGONUCLEOTIDE III, FIGURE 8

PCT-US92-10792-3

Query Match 4.6%; Score 15; DB 5; Length 30;
Best Local Similarity 100.0%; Pred. No. 7.8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 212 ttgtttaaatttg 226
|||||
Db 5 TTTGTTAAAAATTG 19

RESULT 6
US-08-256-261-29/C
Sequence 29, Application US/08256261
Patent No. 5801037
GENERAL INFORMATION:
APPLICANT: Behnke, Detlef
APPLICANT: Schlotz, Bernhard
APPLICANT: Albrecht, Sybille
APPLICANT: Albrecht, Sybille
APPLICANT: G hrs, Karl-Heinz
TITLE OF INVENTION: Expression of signal-peptide-free
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Neave
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/256,261
FILING DATE:
CLASSIFICATION: 435
INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 32 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "synthetic DNA from
DESCRIPTION: oligonucleotide synthesis"
US-08-256-261-29

Query Match 4.6%; Score 15; DB 1; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.7e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 154 aaaggaataataa 168
|||||
Db 30 AAAGGAATAATAA 16

RESULT 7
US-08-852-299-29/C
Sequence 29, Application US/08852299
Patent No. 6010897
GENERAL INFORMATION:
APPLICANT: Behnke, Detlef
APPLICANT: Schlotz, Bernhard
APPLICANT: Albrecht, Sybille
APPLICANT: G hrs, Karl-Heinz
APPLICANT: Hartmann, Manfred
TITLE OF INVENTION: Expression of signal-peptide-free

TITLE OF INVENTION: staphylokinases
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Neave
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/852,299
FILING DATE: 17-MAY-1997
CLASSIFICATION: 435
PRIOR APPLICATION NUMBER: 08/256,261
FILING DATE:
INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 32 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "synthetic DNA from
DESCRIPTION: oligonucleotide synthesis"
US-08-852-299-29

Query Match 4.6%; Score 15; DB 3; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.7e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 154 aaaggaataataaa 168
|||||
DB 30 AAAGGAATAATAAA 16

RESULT 8
US-09-205-204-20/c
Sequence 20, Application US/09205204
Patent No. 5958772
GENERAL INFORMATION:
APPLICANT: C. Frank Bennett
APPLICANT: Elizabeth J. Ackermann
TITLE OF INVENTION: ANTISENSE MODULATION OF CELLULAR INHIBITOR OF APOPTOSIS-1 EXPRESS
FILE REFERENCE: RTS-0020
CURRENT APPLICATION NUMBER: US/09/205,204
CURRENT FILING DATE: 1998-12-03
NUMBER OF SEQ ID NOS: 47
SEQ ID NO 20
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Antisense oligonucleotide
US-09-205-204-20

Query Match 4.3%; Score 14; DB 2; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 160 aaataataaaa 173
|||||
DB 18 AAATAATAATAAA 5

RESULT 9
US-08-816-977-12/c
Sequence 12, Application US/08816977
Patent No. 6080400
GENERAL INFORMATION:
APPLICANT: Williams, James A.
APPLICANT: Byrne, Lisa M.
APPLICANT: Pugh, Charles S.G.
TITLE OF INVENTION: Prevention And Treatment Of
Verotoxin-Induced Disease
NUMBER OF SEQUENCES: 49
CORRESPONDENCE ADDRESS:
ADDRESSEE: Medlen & Carroll, LLP
STREET: 220 Montgomery Street, Suite 2200
CITY: San Francisco
STATE: California
COUNTRY: United States of America
ZIP: 94104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/816,977
FILING DATE: 13-MAR-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: MacKnight, Kamlin T.
REGISTRATION NUMBER: 38,230
REFERENCE/DOCKET NUMBER: OPMD-02450
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 705-8410
TELEFAX: (415) 397-8338
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 29 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-816-977-12

Query Match 4.3%; Score 14; DB 3; Length 29;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataataattt 202
|||||
DB 22 AAATAATATTTT 9

RESULT 10
US-08-629-001A-31
Sequence 31, Application US/08629001A
Patent No. 5858661
GENERAL INFORMATION:
APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
NUMBER OF SEQUENCES: 139
CORRESPONDENCE ADDRESS:
ADDRESSEE: Kohn & Associates
STREET: 30500 No. 5858661Western Hwy.
CITY: Farmington Hills
STATE: Michigan
COUNTRY: US
ZIP: 48334
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,001A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Kohn, Kenneth I.
REGISTRATION NUMBER: 30,955
REFERENCE/DOCKET NUMBER: 2290,00032
TELECOMMUNICATION INFORMATION:
TELEPHONE: (810) 539-5050
TELEFAX: (810) 539-5055
INFORMATION FOR SEQ ID NO: 31:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-629-001A-31

Query Match 4.3%; Score 14; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttattta 209
|||||
Db 14 ttattttattttta 27

RESULT 11
US-08-642-274D-110
Sequence 110, Application US/08642274D
Patent No. 6200749
GENERAL INFORMATION:
APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
TITLE OF INVENTION: SCREEN FOR A PARTIAL A-T PHENOTYPE
FILE REFERENCE: 229000033
CURRENT APPLICATION NUMBER: US/08/642,274D
CURRENT FILING DATE: 1996-05-03
NUMBER OF SEQ ID NOS: 220
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 110
LENGTH: 30
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence:intronic
US-08-642-274D-110

Query Match 4.3%; Score 14; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttattta 209
|||||
Db 14 ttattttattttta 27

RESULT 12
US-08-629-600-16/c
Sequence 16, Application US/08629600
Patent No. 5783196
GENERAL INFORMATION:
APPLICANT: NORIEGA, Fernando
APPLICANT: LEVINE, Myron M.
TITLE OF INVENTION: GOA MUTANTS OF SHIGELLA
TITLE OF INVENTION: AND VACCINES CONTAINING THE SAME
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:

ADDRESSEE: SUGHRUE, MIOM, ZINN, MACPEAK & SEAS
STREET: 2100 Pennsylvania Avenue, N.W., Suite 800
CITY: Washington, D.C.
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20037

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,600
FILING DATE: 9-APR-1996

CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: KIT, Gordon
REGISTRATION NUMBER: 30,764
REFERENCE/DOCKET NUMBER: A-6765
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 293-7060
TELEFAX: (202) 293-7860
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
HYPOTHETICAL: NO

US-08-629-600-16

Query Match 4.3%; Score 14; DB 1; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttt 202
|||||
Db 27 AAATAATTATTTT 14

RESULT 13
US-08-403-853-8
Sequence 8, Application US/08403853
Patent No. 5844094
GENERAL INFORMATION:
APPLICANT: HUDSON, Peter J.
APPLICANT: LAH, Maria
APPLICANT: KORRT, Alex A.
APPLICANT: IRVING, Robert A.
APPLICANT: ATWELL, John L.
APPLICANT: MALBY, Robyn L.
APPLICANT: POWER, Barbara E.
APPLICANT: COLEMAN, Peter M.
TITLE OF INVENTION: TARGET BINDING POLYPEPTIDE
NUMBER OF SEQUENCES: 25
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/403,853
FILING DATE: 30-MAY-1995
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/AU93/00491
FILING DATE: 24-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: AU PL 4973
FILING DATE: 25-SEP-1992
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16786/189/CHAC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-403-853-8

Query Match 4.3%; Score 14; DB 2; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 83 aaattataattat 96
|||||
Db 1 AAATTATATATAT 14

RESULT 14

US-08-847-844A-113
Sequence 113, Application US/08847844A
Patent No. 6150160
GENERAL INFORMATION:
APPLICANT: KAZAZIAN JR., HAIG H.
APPLICANT: BOEKE, JEFF D.
APPLICANT: MORAN, JOHN V.
APPLICANT: DOMBROSKI, BETH A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS OF USE OF
TITLE OF INVENTION: MAMMALIAN RETROTRANSPOSONS
NUMBER OF SEQUENCES: 137
CORRESPONDENCE ADDRESS:
ADDRESSEE: PATITCH SCHWARZE JACOBS & NADEL, P.C.
STREET: ONE COMMERCE SQUARE, 2005 MARKET STREET, 22ND FL.
CITY: PHILADELPHIA
STATE: PA
COUNTRY: U.S.A.
ZIP: 19103-7086
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/847,844A
FILING DATE: 28-APR-1997
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/749,805
FILING DATE: 16-NOV-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/006,831
FILING DATE: 16-NOV-1995
ATTORNEY/AGENT INFORMATION:
NAME: DOYLE LEARY Ph.D., KATHRYN
REGISTRATION NUMBER: 36,317
REFERENCE/DOCKET NUMBER: 9596-2302
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-567-2020
TELEFAX: 215-567-2991

INFORMATION FOR SEQ ID NO: 113:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-847-844A-113

Query Match 4.0%; Score 13; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.3e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 150 ttaaaaggagaaa 162
|||||
Db 5 TTAAAGAGGAGAAA 17

RESULT 15

US-08-686-968C-13
Sequence 13, Application US/08686968C
Patent No. 6221361
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
APPLICANT: Junker, David E.
TITLE OF INVENTION: Recombinant Swinepox Virus
FILE REFERENCE: 39119-H/OML
CURRENT APPLICATION NUMBER: US/08/686,968C
CURRENT FILING DATE: 1996-07-25
NUMBER OF SEQ ID NOS: 231
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 13
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: primer
US-08-686-968C-13

Query Match 4.0%; Score 13; DB 4; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.3e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aattgaattgttaa 243
|||||
Db 5 aattgaattgttaa 17

RESULT 16

US-09-288-461-79
Sequence 79, Application US/09288461
Patent No. 6159694
GENERAL INFORMATION:
APPLICANT: Karras, James G.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of STAT3
FILE REFERENCE: ISPH-0338
CURRENT APPLICATION NUMBER: US/09/288,461
CURRENT FILING DATE: 1999-04-08
NUMBER OF SEQ ID NOS: 107
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 79
LENGTH: 20
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-09-288-461-79

Query Match 4.0%; Score 13; DB 3; Length 20;

Best Local Similarity 100.0%; Pred. No. 5.2e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 173 attagataaag 185
|||||

Db 2 attagataaag 14

RESULT 17

US-08-691-045-61/c
Sequence 61, Application US/08691045
Patent No. 6015664
GENERAL INFORMATION:
APPLICANT: Henrickson, Kelly J.
TITLE OF INVENTION: VIRUS ASSAY METHOD
NUMBER OF SEQUENCES: 65
CORRESPONDENCE ADDRESS:
ADDRESSEE: Quarles & Brady
STREET: 411 East Wisconsin Avenue
CITY: Milwaukee
STATE: Wisconsin
COUNTRY: U.S.A.
ZIP: 53202-4497
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/691,045
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 650053.91037
TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5000
TELEFAX: (414) 271-3552
INFORMATION FOR SEQ ID NO: 61:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Oligonucleotide
US-08-691-045-61

Query Match 4.0%; Score 13; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 94 tatatgataagta 106
|||||

Db 17 TATATGATAGTA 5

RESULT 18

US-08-672-215-1/c
Sequence 1, Application US/08672215
Patent No. 6020121
GENERAL INFORMATION:
APPLICANT: Ying Bao, Amy Boggs, Pamela R. Contag,
APPLICANT: Nancy A. Federspiel, Alan Herbert,
APPLICANT: Scott J. Hecker, Francois Malouin
TITLE OF INVENTION: INHIBITORS OF REGULATORY PATHWAYS
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700

CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066

COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" diskette, 1.44 Mb
COMPUTER: IBM compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq for Windows 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/672,215
FILING DATE: June 25, 1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/004,626
FILING DATE: September 29, 1995
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Walburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 219/158
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-672-215-1

Query Match 4.0%; Score 13; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.1e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 294 tgaattttatg 306
|||||

Db 20 TGAATTTTATG 8

RESULT 19

US-08-120-827-64
Sequence 64, Application US/08120827
Patent No. 5525495
GENERAL INFORMATION:
APPLICANT: KEENE, JACK D.
APPLICANT: KING, PETER H.
APPLICANT: LEVINE, TODD
TITLE OF INVENTION: METHODS AND COMPOSITIONS USEFUL IN THE
RECOGNITION, BINDING AND EXPRESSION OF RIBONUCLEIC ACIDS
INVOLVED IN CELL GROWTH, NEOPLASIA AND IMMUNOREGULATION
NUMBER OF SEQUENCES: 101
CORRESPONDENCE ADDRESS:
ADDRESSEE: OBLON, SPIVAK, MCLELLAND, MAIER & NEUSTADT,
ADDRESS: P. C.
STREET: 1755 Jefferson Davis Highway, Fourth Floor
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/120,827
FILING DATE: 15-SEP-1993

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;
;   TYPE: nucleic acid
STRANDEDNESS: unknown
;
```

US-08-629-001A-79

; TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
 ; TITLE OF INVENTION: SCREEN FOR A PARTIAL A-T PHENOTYPE
 ; FILE REFERENCE: 729000033

; CURRENT APPLICATION NUMBER: US/08/642,274D
; CURRENT FILING DATE: 1996-05-03
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 158
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:intronic
; OTHER INFORMATION: sequence
US-08-642-274D-158

Query Match 4.0%; Score 13; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 272 aaaaattatttc 284
|||
Db 15 aaaaattatttc 27

RESULT 23
US-08-330-638D-5/c
; Sequence 5, Application US/08330638D
; Patent No. 5731425
; GENERAL INFORMATION:
; APPLICANT: Brizard, Billy
; APPLICANT: Bianca, Darlene
; APPLICANT: Chubert, Richard
; APPLICANT: Vizard, Douglas
; APPLICANT: Hopp, Thomas
; TITLE OF INVENTION: POLYPEPTIDE SURFACE
; TITLE OF INVENTION: MARKER FOR CELLS
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Eastman Kodak Company,
; ADDRESSEE: Patent Legal Staff
; STREET: 343 State Street
; CITY: Rochester
; STATE: New York
; COUNTRY: U.S.A.
; ZIP: 14650-2201
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch,
; MEDIUM TYPE: 1.44 MB storage, (Hewlett Packard)
; COMPUTER: HP Vectra
; OPERATING SYSTEM: MS-DOS Version 6.0
; SOFTWARE: WORD FOR WINDOWS
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/330,638D
; FILING DATE: 28 OCT 1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA: NONE
; ATTORNEY/AGENT INFORMATION:
; NAME: Kiernan, Anne B
; REGISTRATION NUMBER: 36,566
; REFERENCE/DOCKET NUMBER: 71255
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (716) 588-2405
; TELEFAX: (716) 477-4646
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 BASES
; TYPE: NUCLEIC ACID
; STRANDEDNESS: SINGLE
; TOPOLOGY: LINEAR
; MOLECULE TYPE: SYNTHETIC OLIGONUCLEOTIDE
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE: SYNTHETICALLY PREPARED
; IMMEDIATE SOURCE: SYNTHETICALLY PREPARED

; PUBLICATION INFORMATION: NONE
; US-08-330-638D-5

Query Match 4.0%; Score 13; DB 1; Length 31;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtctcggggg 136
|||||
Db 28 GTGTCTCGGGGG 16

RESULT 24
US-08-906-746A-5/c
; Sequence 5, Application US/08906746A
; Patent No. 5945292
; GENERAL INFORMATION:
; APPLICANT: Brizard, Billy L.
; APPLICANT: Bianca, Darlene W.
; APPLICANT: Chubert, Richard G.
; APPLICANT: Vizard, Douglas L.
; APPLICANT: Hopp, Thomas P.
; TITLE OF INVENTION: Method of Identifying Cells with
; TITLE OF INVENTION: Polypeptide Surface Marker
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Senniger, Powers, Leavitt & Roedel
; STREET: One Metropolitan Square- 16th floor
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/906,746A
; FILING DATE: 06-AUG-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Stone, Paul A.
; REGISTRATION NUMBER: 38,628
; REFERENCE/DOCKET NUMBER: SGM 6874
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 314-231-5400
; TELEFAX: 314-231-4342
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; US-08-906-746A-5

Query Match 4.0%; Score 13; DB 2; Length 31;
Best Local Similarity 100.0%; Pred. No. 4.9e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtctcggggg 136
|||||
Db 28 GTGTCTCGGGGG 16

RESULT 25
US-08-247-809A-14/c
; Sequence 14, Application US/08247809A

Patent No. 5569823
GENERAL INFORMATION:
APPLICANT: Peter H. Schreier; Klaus Stenzel; Gunter Adam;
APPLICANT: Edgar Maiss
TITLE OF INVENTION: DEOXYRIBONUCLEIC ACIDS
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: SPRUNG HORN KRAMER & WOODS
STREET: 660 White Plains Road
CITY: Tarrytown
STATE: New York
COUNTRY: U.S.A.
ZIP: 10591-5144
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 2.0 MB
MEDIUM TYPE: storage
COMPUTER: NEC Powermate 1 Plus
OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/247,809A
FILING DATE: May 23, 1994
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P 43 178 45.6 (Germany)
FILING DATE: May 28, 1993
ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: Bayer 9049-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-247-809A-14

Query Match 4.0%; Score 13; DB 1; Length 36;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 156 aggaataataaa 168
|||||
Db 24 AGGAATAATATAA 12

RESULT 26
US-08-711-728-14/c
Sequence 14, Application US/08711728
Patent No. 5973135
GENERAL INFORMATION:
APPLICANT: Peter H. Schreier; Klaus Stenzel; Gunter Adam;
APPLICANT: Edgar Maiss
TITLE OF INVENTION: DEOXYRIBONUCLEIC ACIDS
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: SPRUNG HORN KRAMER & WOODS
STREET: 660 White Plains Road
CITY: Tarrytown
STATE: New York
COUNTRY: U.S.A.
ZIP: 10591-5144
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 2.0 MB
MEDIUM TYPE: storage
COMPUTER: NEC Powermate 1 Plus

OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/711,728
FILING DATE: 03-SEPT-1996
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/247,809
FILING DATE: 23-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DE 43178456
FILING DATE: 28-MAY-1993
ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: Bayer 9049.1-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
TELEX:
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-711-728-14

Query Match 4.0%; Score 13; DB 2; Length 36;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 156 aggaataataaa 168
|||||
Db 24 AGGAATAATATAA 12

RESULT 27
US-08-097-554A-45
Sequence 45, Application US/08097554A
Patent No. 5869312
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
APPLICANT: Junker M.S., David E
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 112
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/097,554A
FILING DATE: July 22, 1993
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs

```

: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHETICAL: NO
: ANTI-SENSE: NO
: ORIGINAL SOURCE:
: ORGANISM: Swinepox virus
: STRAIN: Kasza
: INDIVIDUAL ISOLATE: S-SPV-001
: IMMEDIATE SOURCE:
: CLONE: 515-85.1
: POSITION IN GENOME:
: MAP POSITION: -23.2
: UNITS: %G
:
: US-08-097-554A-45

```

```

Query Match
Best Local Similarity 100.0%; Score 13; DB 2; Length 37;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

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QY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

```

```

RESULT 28
US-08-484-575A-10
: Sequence 10, Application US/08484575A
: Patent No. 5925358
: GENERAL INFORMATION:
: APPLICANT: Mark D. Cochran and David E. Junker
: TITLE OF INVENTION: Recombinant Fowlpox viruses and Uses Thereof
: NUMBER OF SEQUENCES: 42
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: John P. White
: STREET: 1185 Avenue of the Americas
: CITY: New York
: STATE: New York
: COUNTRY: USA
: ZIP: 10036
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: OPERATING SYSTEM: IBM PC compatible
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/484,575A
: FILING DATE: 07-JUN-1995
: CLASSIFICATION: 424
: ATTORNEY/AGENT INFORMATION:
: NAME: White Esq, John P
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212)278-0450
: TELEFAX: (212)391-0525
: INFORMATION FOR SEQ ID NO: 10:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 37 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHETICAL: NO
: ANTI-SENSE: NO
:
: US-08-484-575A-10

```

```

Query Match
Best Local Similarity 100.0%; Score 13; DB 2; Length 37;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 231 aatgaattgttaa 243

```

```

Db 10 AATGAATTGTAA 22

```

```

RESULT 29
US-08-477-459-10
: Sequence 10, Application US/08477459
: Patent No. 6001369
: GENERAL INFORMATION:
: APPLICANT: Mark D. Cochran
: TITLE OF INVENTION: Recombinant Fowlpox viruses and Uses
: NUMBER OF SEQUENCES: 20
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: John P. White
: STREET: 1185 Avenue of the Americas
: CITY: New York
: STATE: New York
: COUNTRY: USA
: ZIP: 10036
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: OPERATING SYSTEM: IBM PC compatible
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/477,459
: FILING DATE: 07-JUN-1995
: CLASSIFICATION: 424
: ATTORNEY/AGENT INFORMATION:
: NAME: White Esq, John P
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212) 278-0400
: TELEFAX: (212) 391-0525
: INFORMATION FOR SEQ ID NO: 10:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 37 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHETICAL: NO
: ANTI-SENSE: NO
:
: US-08-477-459-10

```

```

Query Match
Best Local Similarity 100.0%; Score 13; DB 3; Length 37;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

```

```

RESULT 30
US-08-480-640A-45
: Sequence 45, Application US/08480640A
: Patent No. 6033904
: GENERAL INFORMATION:
: APPLICANT: Cochran, Mark D.
: TITLE OF INVENTION: Recombinant Swinepox Virus
: NUMBER OF SEQUENCES: 225
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: John P. White
: STREET: 1185 Avenue of the Americas
: CITY: New York
: STATE: New York
: COUNTRY: USA
: ZIP: 10036
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk

```

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/08/480,640A
APPLICATION NUMBER: US/08/480,640A
FILING DATE: 07-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
REGISTRATION NUMBER: 28,678
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: -23.2
UNITS: %
US-08-480-640A-45

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 231 aattgaattgtaa 243
DB 10 AATTGAATTGTAA 22

RESULT 31
US-08-479-869-10
Sequence 10, Application US/08479869
Patent No. 6123949
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
TITLE OF INVENTION: Recombinant Fowlpox Virus S-PPV-043 and
TITLE OF INVENTION: Uses Thereof
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/479,869
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/024,156
FILING DATE: 26-FEB-1993
ATTORNEY/AGENT INFORMATION:
NAME: White Esq, John P

TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
US-08-479-869-10

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 231 aattgaattgtaa 243
DB 10 AATTGAATTGTAA 22

RESULT 32
US-08-295-802-45
Sequence 45, Application US/08295802
Patent No. 6127163
GENERAL INFORMATION:
APPLICANT: Cochran Ph.D., Mark D
APPLICANT: Junker M.S., David E
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/295,802
FILING DATE: Herewith
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: -23.2

UNITS: %G
US-08-295-802-45

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

RESULT 33
US-08-486-414-10
Sequence 10, Application US/08486414B
Patent No. 6136318
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUSES AND USES THEREOF
FILE REFERENCE: 42771D
CURRENT APPLICATION NUMBER: US/08/486,414B
CURRENT FILING DATE: 1995-06-07
NUMBER OF SEQ ID NOS: 46
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 10
LENGTH: 37
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Linker
US-08-486-414-10

Query Match 4.0%; Score 13; DB 3; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 aatgaattgttaa 22

RESULT 34
US-08-488-237A-45
Sequence 45, Application US/08488237A
Patent No. 6251403
GENERAL INFORMATION:
APPLICANT: Cochran, Mark D.
TITLE OF INVENTION: Recombinant Swinepox Virus
NUMBER OF SEQUENCES: 225
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 1185 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/488,237A
FILING DATE: 07-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: White, John P
REGISTRATION NUMBER: 28,678
TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Swinepox virus
STRAIN: Kasza
INDIVIDUAL ISOLATE: S-SPV-001
IMMEDIATE SOURCE:
CLONE: 515-85.1
POSITION IN GENOME:
MAP POSITION: -23.2
UNITS: %G
US-08-488-237A-45

Query Match 4.0%; Score 13; DB 4; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
Db 10 AATGAATTGTAA 22

RESULT 35
PCT-US94-01826A-10
Sequence 10, Application PC/TUS9401826A
GENERAL INFORMATION:
APPLICANT: Syntro Corporation, et al.
TITLE OF INVENTION: Recombinant Fowlpox Virus S-FPV-043 and Uses Thereof
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: John P. White
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/01826A
FILING DATE: 28-FEB-1994
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: White Esq, John P
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)977-9550
TELEFAX: (212)664-0525
TELEX: 422523
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 37 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
PCT-US94-01826A-10

Query Match 4.0%; Score 13; DB 5; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
|||||
Db 10 AATGAATTGTAA 22

RESULT 36

PCT-US94-02252A-10
; Sequence 10, Application PC/TUS9402252A
; GENERAL INFORMATION:
; APPLICANT: Syntro Corporation, et al.
; TITLE OF INVENTION: Recombinant Fowlpox viruses and uses thereof
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John P. White
; STREET: 30 Rockefeller Plaza
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10112
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/02252A
; FILING DATE: 28-FEB-1994
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: White Esq, John P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212)977-9550
; TELEFAX: (212)664-0525
; TELEX: 422523
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 37 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; PCT-US94-02252A-10

Query Match 4.0%; Score 13; DB 5; Length 37;
Best Local Similarity 100.0%; Pred. No. 4.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgttaa 243
|||||
Db 10 AATGAATTGTAA 22

RESULT 37

PCT-US96-00547-40/C
; Sequence 40, Application PC/TUS9600547
; GENERAL INFORMATION:
; APPLICANT: Virogenetics Corporation
; TITLE OF INVENTION: RECOMBINANT POXVIRUS-HTLV, COMPOSITIONS
; NUMBER OF SEQUENCES: 56
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford, P.C.
; STREET: 530 Fifth Avenue, 25th Floor
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.

ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/00547
; FILING DATE: 12-JAN-1996
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/372,664
; FILING DATE: 13-JAN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Frommer, William S.
; REGISTRATION NUMBER: 25,506
; REFERENCE/DOCKET NUMBER: 454310-2621
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 840-3333
; TELEFAX: (212) 840-0712
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 38 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; PCT-US96-00547-40

Query Match 4.0%; Score 13; DB 5; Length 38;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 293 ctgtaattttat 305
|||||
Db 25 CTGTAATTTTAT 13

RESULT 38
US-08-105-483-168/C
; Sequence 168, Application US/08105483
; Patent No. 5494807
; GENERAL INFORMATION:
; APPLICANT: Paolelli, Enzo
; TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
; NUMBER OF SEQUENCES: 462
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford
; ADDRESSEE: C/O William S. Frommer
; STREET: 530 Fifth Avenue
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/105,483
; FILING DATE: 12-AUG-1993
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/847,951
; FILING DATE: 06-MAR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Frommer, William S.
; REGISTRATION NUMBER: 25,506
; REFERENCE/DOCKET NUMBER: 454310-2400
; TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 840-3333
 TELEFAX: (212) 840-0712
 INFORMATION FOR SEQ ID NO: 168:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 39 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-105-483-168

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 293 ctgtaattttat 305
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 39
 US-08-709-209-168/c
 Sequence 168, Application US/08709209
 Patent No. 5762938
 GENERAL INFORMATION:
 APPLICANT: Paoletti, Enzo
 TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
 TITLE OF INVENTION: STRAIN
 NUMBER OF SEQUENCES: 462
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Curtis, Morris & Safford
 ADDRESSEE: c/o William S. Frommer
 STREET: 530 Fifth Avenue
 CITY: New York
 STATE: NY
 COUNTRY: USA
 ZIP: 10036
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/709,209
 FILING DATE: 21-AUG-1996
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/105,483
 FILING DATE: 12-AUG-1993
 APPLICATION NUMBER: US 07/847,951
 FILING DATE: 06-MAR-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: Frommer, William S.
 REGISTRATION NUMBER: 25,506
 REFERENCE/DOCKET NUMBER: 454310-2400
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212) 840-3333
 TELEFAX: (212) 840-0712
 INFORMATION FOR SEQ ID NO: 168:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 39 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-709-209-168

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 293 ctgtaattttat 305
 |||||||

Db 25 CTGTAATTTTAT 13

RESULT 40
 US-08-303-275-56/c
 Sequence 56, Application US/08303275
 Patent No. 5766598
 GENERAL INFORMATION:
 APPLICANT: Paoletti, Enzo
 APPLICANT: Tartaglia, James
 APPLICANT: Cox, William I.
 TITLE OF INVENTION: IMMUNODEFICIENCY VIRUS RECOMBINANT
 TITLE OF INVENTION: POXVIRUS VACCINE
 NUMBER OF SEQUENCES: 205
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Curtis, Morris & Safford
 ADDRESSEE: c/o William S. Frommer
 STREET: 530 Fifth Avenue
 CITY: New York
 STATE: New York
 COUNTRY: USA
 ZIP: 10036
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/303,275
 FILING DATE:
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 07/897,382
 FILING DATE: 11-JUN-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: Frommer, William S.
 REGISTRATION NUMBER: 25,506
 REFERENCE/DOCKET NUMBER: 454310-2420
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212) 840-3333
 TELEFAX: (212) 840-0712
 INFORMATION FOR SEQ ID NO: 56:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 39 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-303-275-56

Query Match 4.0%; Score 13; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 4.7e+03;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 293 ctgtaattttat 305
 |||||||
 Db 25 CTGTAATTTTAT 13

RESULT 41
 US-08-458-101-168/c
 Sequence 168, Application US/08458101
 Patent No. 5766599
 GENERAL INFORMATION:
 APPLICANT: Paoletti, Enzo
 APPLICANT: Perkus, Marion E.
 APPLICANT: Taylor, Jill
 APPLICANT: Tartaglia, James
 APPLICANT: No. 5766599ton, Elizabeth K.
 APPLICANT: Riviere, Michel
 APPLICANT: de Taisne, Charles
 APPLICANT: Limbach, Keith J.
 APPLICANT: Johnson, Gerard P.

APPLICANT: Pincus, Steven E.
APPLICANT: Cox, William I.
APPLICANT: Audonnet, Jean-Christophe Francis
APPLICANT: Getlig, Russell Robert
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
TITLE OF INVENTION: STRAIN
NUMBER OF SEQUENCES: 467
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Curtis, Morris & Safford
ADDRESSEE: C/O William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/458,101
FILING DATE: 01-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2740
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-458-101-168

Query Match 4.0%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 293 ctgtaattttat 305
|||||
DB 25 CTGTAATTTTAT 13

RESULT 42
US-09-028-361A-19
Sequence 19, Application US/09028361A
Patent No. 5962296
GENERAL INFORMATION:
APPLICANT: ETWILLER, LAURENCE
APPLICANT: XU, SHUANG-YONG
TITLE OF INVENTION: METHOD FOR CLONING AND
TITLE OF INVENTION: PRODUCING THERMICROBIUM ROSEUM DNA
TITLE OF INVENTION: POLYMERASE I IN E. COLI
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: New England Biolabs, Inc.
STREET: 32 Tozer Road
CITY: Beverly
STATE: MA
COUNTRY: USA
ZIP: 01915
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/028,361A
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Williams, Gregory D.
REGISTRATION NUMBER: 30901
REFERENCE/DOCKET NUMBER: NEB-134
TELECOMMUNICATION INFORMATION:
TELEPHONE: 978-927-5054
TELEFAX: 978-927-1705
TELEX:
INFORMATION FOR SEQ ID NO: 19:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-09-028-361A-19

Query Match 4.0%; Score 13; DB 2; Length 39;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 62 atataaattat 74
|||||
DB 12 ATATATTAATAT 24

RESULT 43
US-08-199-507B-39/C
Sequence 39, Application US/08199507B
Patent No. 5472841
GENERAL INFORMATION:
APPLICANT: JAYASENA, S. AND GOLD, L.
TITLE OF INVENTION: NUCLEIC ACID LIGAND INHIBITORS
NUMBER OF SEQUENCES: 59
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Swanson & Bratschun, L.L.C.
STREET: 8400 E. Prentice Avenue, Suite 200
CITY: Englewood
STATE: Colorado
COUNTRY: USA
ZIP: 80111
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 Inch, 360 KB storage
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/199,507B
FILING DATE: 22 FEBRUARY 1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX13
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 39:
SEQUENCE CHARACTERISTICS:

LENGTH: 40 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-199-507B-39

Query Match 4.0%; Score 13; DB 1; Length 40;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 157 ggaataataat 169
|||||
DB 22 GGAATAATAAT 10

RESULT 44
US-08-441-828-39/C
Sequence 39, Application US/08441828
Patent No. 5734034
GENERAL INFORMATION:
APPLICANT: JAYASENA, S. AND GOLD, L.
TITLE OF INVENTION: NUCLEIC ACID LIGAND INHIBITORS
TITLE OF INVENTION: OF HUMAN NEUTROPHIL ELASTASE
NUMBER OF SEQUENCES: 59
CORRESPONDENCE ADDRESS:
ADDRESSEE: Swanson & Bratschun, L.L.C.
STREET: 8400 E. Prentice Avenue, Suite 200
CITY: Englewood
STATE: Colorado
COUNTRY: USA
ZIP: 80111
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 MB
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/441,828
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/199,507
FILING DATE: 22 FEBRUARY 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX13
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 39:
SEQUENCE CHARACTERISTICS:
LENGTH: 40 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-441-828-39

Query Match 4.0%; Score 13; DB 1; Length 40;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 157 ggaataataat 169
|||||

DB 22 GGAATAATAAT 10

RESULT 45
US-08-930-503A-11
Sequence 11, Application US/08930503A
Patent No. 6132731
GENERAL INFORMATION:
APPLICANT:
TITLE OF INVENTION: MORINE LEUKEMIA VIRUS VECTORS (AS AMENDED)
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: WENDEROTH, LIND & PONACK, L.L.P.
STREET: 2033 K Street, N.W. - Suite 800
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20006
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/930,503A
FILING DATE: October 8, 1997
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9506782.3
FILING DATE: 01-APR-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/GB96/00776
FILING DATE: 01-APR-1996
ATTORNEY/AGENT INFORMATION:
NAME: Warren M Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-721-8200
TELEFAX: 202-721-8250
TELEX:
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 41 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-930-503A-11

Query Match 4.0%; Score 13; DB 3; Length 41;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 251 ttccaggggggaat 263
|||||
DB 2 TTCAGGGGGGAAT 14

Search completed: January 24, 2002, 03:24:24
Job time: 3711 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:19:03 ; Search time 1494.92 Seconds

(without alignments)
3608.606 Million cell updates/sec

Title: US-09-531-438-3

Sequence: 1 atttggatattcttaattt.....ttcatgtttttatttgc 327

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 1472140 seqs, 8248589755 residues

Word size: 0

Total number of hits satisfying chosen parameters: 541028

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

GenEmbl:*

- 1: gb_da:*
- 2: gb_bg:*
- 3: gb_in:*
- 4: gb_ov:*
- 5: gb_ov:*
- 6: gb_pat:*
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- 8: gb_pl:*
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- 11: gb_sts:*
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- 13: gb_un:*
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- 27: em_sy:*
- 28: em_un:*
- 29: em_vl:*
- 30: em_htgo_hum:*
- 31: em_htgo_inv:*
- 32: em_htgo_rod:*
- 33: em_htg_hum:*
- 34: em_htg_inv:*
- 35: em_htg_rod:*
- 36: em_htg_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	4.9	29	6	AR099868
2	16	4.9	32	6	E27913
3	16	4.9	36	3	CEANONVPR
4	16	4.9	36	6	AR019036
5	15	4.6	27	6	AR014030
6	15	4.6	27	6	I21980
7	15	4.6	27	6	AR037189
8	14	4.3	29	6	AR076353
9	14	4.3	30	6	AR099868
10	14	4.3	30	6	AR028182
11	14	4.3	30	6	AR138585
12	14	4.3	32	6	E27913
13	14	4.3	36	6	AR019036
14	14	4.3	37	6	AR063204
15	14	4.3	45	6	AX049973
16	14	4.3	45	6	AX049974
17	14	4.3	45	6	AX099855
18	14	4.3	45	6	AX099856
19	14	4.3	45	6	AX137975
20	14	4.3	45	6	AX137976
21	14	4.3	50	6	AX159856
22	13	4.0	20	6	AR146953
23	13	4.0	20	6	AR121058
24	13	4.0	20	6	AX076045
25	13	4.0	23	6	A97479
26	13	4.0	24	6	AX093544
27	13	4.0	24	6	AX164353
28	13	4.0	24	12	AB069100
29	13	4.0	25	6	AX042574
30	13	4.0	25	6	AX043268
31	13	4.0	26	6	AX039624
32	13	4.0	26	6	AX039654
33	13	4.0	28	6	AR014028
34	13	4.0	28	6	I21978
35	13	4.0	29	6	AX012366
36	13	4.0	29	6	E59972
37	13	4.0	30	6	AR028230
38	13	4.0	30	6	AR138633
39	13	4.0	30	6	AX063379
40	13	4.0	31	6	I95122
41	13	4.0	33	5	XELARSE59
42	13	4.0	36	6	AA1027
43	13	4.0	36	6	AR082586
44	13	4.0	36	6	AX167671
45	13	4.0	36	6	I28261

ALIGNMENTS

RESULT 1

AR099868 LOCUS AR099868 29 bp DNA

DEFINITION Sequence 12 from patent US 6080400. PAT 14-FEB-2001

ACCESSION AR099868

VERSION AR099868.1 GI:12810316

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 29)

AUTHORS Williams,J.A. and Byrne,L.Marie.

TITLE Compositions for the prevention and treatment of verotoxin-induced disease

JOURNAL Patent: US 6080400-A 12 27-JUN-2000;

FEATURES Location/Qualifiers

source 1..29

BASE COUNT 11 a 2 c 5 g 11 t

BASE COUNT 11 a 2 c 5 g 11 t

Query Match 4.9%; Score 16; DB 6; Length 29;
 Best Local Similarity 100.0%; Pred. No. 9.2e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattattttta 204
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 DB 9 AAAATATTATTATT 24

RESULT 2

E27913 32 bp DNA PAT 07-FEB-2001
 LOCUS E27913
 DEFINITION Method for detecting foreign DNA fragment insert in Vero toxin gene.

ACCESSION E27913
 VERSION E27913.1 GI:13020766
 KEYWORDS JP 1999243996-A/3.
 SOURCE unidentified.
 ORGANISM unidentified.

REFERENCE 1 (bases 1 to 32)
 AUTHORS Masahiro, K., Y. N. N. and Kawamura, K. S.
 TITLE Method for detecting foreign DNA fragment insert in Vero toxin gene
 JOURNAL Patent: JP 1999243996-A 3 14-SEP-1999;

COMMENT

OS Unidentified
 PN JP 1999243996-A/3
 PD 14-SEP-1999
 PE 27-FEB-1998 JP 1998047677
 PR

PI MASAIRO KUSUMOTO, YOSHIAKI NISHIYA, YOSHIIHISA KANAMURA, PI
 KUNIHITO SHINAGAWA
 PC C1201/68,C12N15/09/(C12N15/09,C12R1:185),C12N15/00,
 (C12N15/00, PC C12R1:185)
 CC Strandedness: Both;
 CC Topology: Linear;
 FH key
 FT source
 FT Location/Qualifiers
 1.32 /organism="unidentified".
 Location/Qualifiers

BASE COUNT 12 a 2 c 4 g 14 t
 ORIGIN

Query Match 4.9%; Score 16; DB 6; Length 32;
 Best Local Similarity 100.0%; Pred. No. 9.2e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattattttta 204
 |||||||
 DB 4 AAAATATTATTATT 19

RESULT 3
 CEANONYFR 36 bp DNA INV 06-MAY-1997
 LOCUS CEANONYFR/c
 DEFINITION C elegans DNA fragment with rearrangement site.
 ACCESSION X97532
 VERSION X97532.1 GI:1296496
 KEYWORDS

SOURCE Caenorhabditis elegans.
 ORGANISM Caenorhabditis elegans.

REFERENCE 1 (bases 1 to 36)
 AUTHORS Wicky, C., Villeneuve, A. M., Lauper, N., Codourey, L., Tobler, H. and Muller, F.

Telomeric repeats (TTAGGC)n are sufficient for chromosome capping
 TITLE

JOURNAL function in Caenorhabditis elegans
 Proc. Natl. Acad. Sci. U.S.A. 93 (17), 8983-8988 (1996)
 MEDLINE 96392352
 REFERENCE 2 (bases 1 to 36)
 AUTHORS Wicky, C.
 TITLE Direct Submission
 JOURNAL Submitted (24-APR-1996) C. Wicky, University of British Columbia,
 Medical Genetics, 6174 University Boulevard, Vancouver, B.C. V6T
 123, CANADA

FEATURES

source

1.36 /organism="Caenorhabditis elegans"
 /db_xref="taxon:6239"
 /clone="Bp1"

misc_feature 1.36 /note="sequence with rearrangement site"
 BASE COUNT 9 a 5 c 4 g 18 t
 ORIGIN

Query Match 4.9%; Score 16; DB 3; Length 36;
 Best Local Similarity 100.0%; Pred. No. 9.1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 169 taaatttagataaa 184
 |||||||
 DB 23 TAAATTTAGATRAAA 8

RESULT 4

ARO19036 36 bp DNA PAT 05-DEC-1998
 LOCUS ARO19036
 DEFINITION Sequence 16 from patent US 5783196.
 ACCESSION ARO19036
 VERSION ARO19036.1 GI:3974150
 KEYWORDS

SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 36)
 AUTHORS Noriega, F. R. and Levine, M. M.
 TITLE Gua mutants of shigella spp. and vaccines containing the same
 JOURNAL Patent: US 5783196-A 16 21-JUL-1998;
 FEATURES Location/Qualifiers
 source 1.36 /organism="unknown"

BASE COUNT 11 a 3 c 10 g 12 t
 ORIGIN

Query Match 4.9%; Score 16; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 9.1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattattttta 204
 |||||||
 DB 14 AAAATATTATTATT 29

RESULT 5

ARO14030 27 bp DNA PAT 05-DEC-1998
 LOCUS ARO14030
 DEFINITION Sequence 66 from patent US 5773246.
 ACCESSION ARO14030
 VERSION ARO14030.1 GI:3971484
 KEYWORDS

SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 27)
 AUTHORS Keene, J. D., Levine, T. and Gao, F.
 TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia

JOURNAL and Immunoregulation
Patent: US 5773246-A 66 30-JUN-1998;
FEATURES Location/Qualifiers
SOURCE 1..27
BASE COUNT 7 a 2 c 2 g 16 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 197 tattttattttaaa 211
Db 12 TATTTTATTTTAA 26

RESULT 6
LOCUS 121980 27 bp DNA PAT 07-OCT-1996
DEFINITION Sequence 66 from patent US 5525495.
ACCESSION 121980
VERSION 121980.1 GI:1602334
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 27)
AUTHORS Keene,J.D., Levine,T. and Gao,F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5525495-A 66 11-JUN-1996;
FEATURES Location/Qualifiers
SOURCE 1..27
BASE COUNT 7 a 2 c 2 g 16 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 197 tattttattttaaa 211
Db 12 TATTTTATTTTAA 26

RESULT 7
LOCUS AR037189 32 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 29 from patent US 5801037.
ACCESSION AR037189
VERSION AR037189.1 GI:5955045
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 32)
AUTHORS Behnke,D., Scholtz,B., Albrecht,S., Guhrs,K. and Hartmann,M.
TITLE Expression of signal peptide-free scapapylolkinases
JOURNAL Patent: US 5801037-A 29 01-SEP-1998;
FEATURES Location/Qualifiers
SOURCE 1..32
BASE COUNT 3 a 8 c 4 g 17 t
ORIGIN

Query Match 4.6%; Score 15; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 2.4e+04;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 154 aaaggaataataa 168
Db 30 AAAGGAATAATAA 16

RESULT 8
LOCUS AR076353 18 bp DNA PAT 30-AUG-2000
DEFINITION Sequence 20 from patent US 5958772.
ACCESSION AR076353
VERSION AR076353.1 GI:10003099
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 18)
AUTHORS Bennett,C.Frank, Ackermann,E.J. and Cowser,L.M.
TITLE Antisense inhibition of cellular inhibitor of apoptosis-1 expression
JOURNAL Patent: US 5958772-A 20 28-SEP-1999;
FEATURES Location/Qualifiers
SOURCE 1..18
BASE COUNT 3 a 2 c 1 g 12 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 6.2e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 160 aaataataataa 173
Db 18 AAATAATAATAA 5

RESULT 9
LOCUS AR099868 29 bp DNA PAT 14-FEB-2001
DEFINITION Sequence 12 from patent US 6080400.
ACCESSION AR099868
VERSION AR099868.1 GI:12810316
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 29)
AUTHORS Williams,J.A. and Byrne,L.Marie.
TITLE Compositions for the prevention and treatment of verotoxin-induced disease
JOURNAL Patent: US 6080400-A 12 27-JUN-2000;
FEATURES Location/Qualifiers
SOURCE 1..29
BASE COUNT 11 a 2 c 5 g 11 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 29;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 189 aaataataattt 202
Db 22 AAATAATAATTTT 9

RESULT 10
LOCUS AR028182 30 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 31 from patent US 5858661.

```

ACCESSION   AR028182
VERSION     AR028182.1  GI:5940155
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 30)
AUTHORS     Shiloh,Y.
TITLE       Ataxia-telangiectasia gene and its genomic organization
JOURNAL     Patent: US 5858661-A 31-12-JAN-1999;
FEATURES
   source    1..30
              Location/Qualifiers
BASE COUNT   11 a      1 c      5 g      13 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY  196 ttattttatttta 209
      |||||||
Db  14 TTAATTTTATTTTA 27

RESULT 11
ARI38585
LOCUS       ARI38585 30 bp DNA PAT 16-JUN-2001
DEFINITION Sequence 110 from patent US 6200749.
ACCESSION   ARI38585
VERSION     ARI38585.1 GI:14480930
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 30)
AUTHORS     Shiloh,Y.
TITLE       Mutated forms of the ataxia-telangiectasia gene and method to
JOURNAL     Patent: US 6200749-A 110 13-MAR-2001;
FEATURES
   source    1..30
              Location/Qualifiers
BASE COUNT   11 a      1 c      5 g      13 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY  196 ttattttatttta 209
      |||||||
Db  14 TTAATTTTATTTTA 27

RESULT 12
E27913/c
LOCUS       E27913 32 bp DNA PAT 07-FEB-2001
DEFINITION Method for detecting foreign DNA fragment insert in Vero toxin
ACCESSION   E27913
VERSION     E27913.1 GI:13020766
KEYWORDS    JP 1999243996-A/3.
SOURCE      unidentified.
ORGANISM    unidentified.
REFERENCE   1 (bases 1 to 32)
AUTHORS     Masahiro,K.Y.N.N. and Kawamura,K.S.
TITLE       Method for detecting foreign DNA fragment insert in Vero toxin gene
JOURNAL     Patent: JP 1999243996-A 3 14-SEP-1999;
            TOYOCO CO LTD

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COMMENT
OS          Unidentified
PN          JP 1999243996-A/3
PD          14-SEP-1999
PF          27-FEB-1998 JP 1998047677
PI          MASAHIRO KUSUMOTO,YOSHIKAKI NISHIYA,YOSHIHISA KAWAMURA, PI
KUNIHICO SHINAGAWA
PC          C1201/68,C12N15/09//((C12N15/09,C12R1:185),C12N15/00,
(C12N15/00, PC C12R1:185)
CC          Strandedness: Both;
CC          Topology: Linear;
C1          Location/Qualifiers
FT          1..32
            source    1..32
              Location/Qualifiers
FEATURES
   source    1..32
              Location/Qualifiers
BASE COUNT   12 a      2 c      4 g      14 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 6.1e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY  189 aaataattatttt 202
      |||||||
Db  17 AAAATATATTTT 4

RESULT 13
AR019036/c
LOCUS       AR019036 36 bp DNA PAT 05-DEC-1998
DEFINITION Sequence 16 from patent US 5783196.
ACCESSION   AR019036
VERSION     AR019036.1 GI:3974150
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 36)
AUTHORS     Noriega,F.R. and Levine,M.M.
TITLE       Gua mutants of shigella sp. and vaccines containing the same
JOURNAL     Patent: US 5783196-A 16 21-JUL-1998;
FEATURES
   source    1..36
              Location/Qualifiers
BASE COUNT   11 a      3 c      10 g      12 t
ORIGIN
Query Match  4.3%; Score 14; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY  189 aaataattatttt 202
      |||||||
Db  27 AAAATATATTTT 14

RESULT 14
AR063204
LOCUS       AR063204 37 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 8 from patent US 5844094.
ACCESSION   AR063204
VERSION     AR063204.1 GI:5990895
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 37)
AUTHORS     Hudson,P.John, Iah,M., Kortt,A.Andrew, Irving,R.Alexander,

```

Atwell,J.Leslie, Malby,R.Louise, Power,B.Elaïne and
Colman,P.Malcolm.
Target binding polypeptide
JOURNAL Patent: US 5844094-A 8 01-DEC-1998;
FEATURES Location/Qualifiers
source 1..37

BASE COUNT 14 a 5 c 0 g 18 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 83 aaattcataattat 96
Db 1 AAATTTATATATAT 14

RESULT 15
LOCUS AX049973 45 bp DNA PAT 12-JAN-2001
DEFINITION Sequence 74 from Patent WO0070071.
ACCESSION AX049973
VERSION AX049973.1 GI:12226350
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Boul,A., Havenga,M.J. and Vogels,R.
TITLE Adenovirus derived gene delivery vehicles comprising at least one
JOURNAL element of adenovirus type 35
Patent: WO 0070071-A 74 23-NOV-2000;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Oligonucleotide TATA-plus"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 21 CTGAAATTTATTA 8

RESULT 16
LOCUS AX049974 45 bp DNA PAT 12-JAN-2001
DEFINITION Sequence 75 from Patent WO0070071.
ACCESSION AX049974
VERSION AX049974.1 GI:12226351
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Boul,A., Havenga,M.J. and Vogels,R.
TITLE Adenovirus derived gene delivery vehicles comprising at least one
JOURNAL element of adenovirus type 35
Patent: WO 0070071-A 75 23-NOV-2000;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"

/db_xref="taxon:32630"
/note="Oligonucleotide TATA-min"
BASE COUNT 19 a 7 c 5 g 14 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 29 CTGAAATTTATTA 42

RESULT 17
LOCUS AX099855 45 bp DNA PAT 02-APR-2001
DEFINITION Sequence 4 from Patent WO0120014.
ACCESSION AX099855
VERSION AX099855.1 GI:13538881
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Schouten,G.J., Vogels,R. and Opstelten,D.J.
TITLE Modified adenoviral vectors for use in gene therapy
JOURNAL Patent: WO 0120014-A 4 22-MAR-2001;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
primer_bind 1..45
/note="Primer TATApus"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 79 ctgaaattataa 92
Db 21 CTGAAATTTATTA 8

RESULT 18
LOCUS AX099856 45 bp DNA PAT 02-APR-2001
DEFINITION Sequence 5 from Patent WO0120014.
ACCESSION AX099856
VERSION AX099856.1 GI:13538882
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.

REFERENCE 1 (bases 1 to 45)
AUTHORS Schouten,G.J., Vogels,R. and Opstelten,D.J.
TITLE Modified adenoviral vectors for use in gene therapy
JOURNAL Patent: WO 0120014-A 5 22-MAR-2001;
Introgene B.V. (NL)

FEATURES
source 1..45
Location/Qualifiers
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
primer_bind 1..45
/note="Primer TATApus"
BASE COUNT 19 a 7 c 5 g 14 t

ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 29 CTGAATAATTATTA 42

RESULT 19

LOCUS AX137975/c 45 bp DNA PAT 30-MAY-2001
DEFINITION Sequence 4 from Patent EP1083229.
ACCESSION AX137975
VERSION AX137975.1 GI:14274070
KEYWORDS

SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 45)
ARTIFICIAL SEQUENCE.

AUTHORS Modified adenoviral vectors for use in gene therapy
TITLE Patent: EP 1083229-A 4 14-MAR-2001;
JOURNAL Introgene B.V. (NL)

FEATURES
source location/Qualifiers
1..45
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer"

primer_bind 1..45
/note="Primer TATApJus/"

BASE COUNT 14 a 5 c 7 g 19 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 21 CTGAATAATTATTA 8

RESULT 20

LOCUS AX137976 45 bp DNA PAT 30-MAY-2001
DEFINITION Sequence 5 from Patent EP1083229.
ACCESSION AX137976
VERSION AX137976.1 GI:14274071
KEYWORDS

SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 45)
ARTIFICIAL SEQUENCE.

AUTHORS Modified adenoviral vectors for use in gene therapy
TITLE Patent: EP 1083229-A 5 14-MAR-2001;
JOURNAL Introgene B.V. (NL)

FEATURES
source location/Qualifiers
1..45
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer"

primer_bind 1..45
/note="Primer TATApJus"

BASE COUNT 19 a 7 c 5 g 14 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92
|||||
DB 29 CTGAATAATTATTA 42

RESULT 21

LOCUS AX159856/c 50 bp DNA PAT 22-JUN-2001
DEFINITION Sequence 3184 from Patent WO0140521.
ACCESSION AX159856
VERSION AX159856.1 GI:14541187
KEYWORDS

SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 50)
AUTHORS Shinkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
methods of use thereof
JOURNAL Patent: WO 0140521-A 3184 07-JUN-2001;
Curagen Corporation (US)

FEATURES
source location/Qualifiers
1..50
/organism="Homo sapiens"
/db_xref="taxon:9606"
misc_feature 25..26
/note="Nucleotide deleted between bases 25 and 26
Accession number cg43063075"

misc_feature 26
/note="2 of 2 allelic variants (3183 is other entry)"

BASE COUNT 10 a 10 c 8 g 22 t
ORIGIN

Query Match 4.3%; Score 14; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 6e+04;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 155 aaggaattataa 168
|||||
DB 36 AAGGAATAATTATA 23

RESULT 22

LOCUS AR146953 18 bp DNA PAT 08-AUG-2001
DEFINITION Sequence 13 from patent US 6221361.
ACCESSION AR146953
VERSION AR146953.1 GI:15110756
KEYWORDS

SOURCE unknown.
ORGANISM unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Cochran,M.D. and Junker,D.E.
TITLE Recombinant swinepox virus
JOURNAL Patent: US 6221361-A 13 24-APR-2001;

FEATURES
source location/Qualifiers
1..18
/organism="unknown"
/db_xref="taxon:32630"
/note="primer"

primer_bind 1..18
/note="Primer TATApJus"

BASE COUNT 8 a 1 c 2 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 231 aatgaattgtaa 243
|||||
Db 5 AATTGAATTGTTAA 17

RESULT 23
LOCUS AR121058 20 bp DNA
DEFINITION Sequence 79 from patent US 6159694.
ACCESSION AR121058
VERSION AR121058.1 GI:14104634
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Kairas,J.G.
TITLE Antisense modulation of stat3 expression
JOURNAL Patent: US 6159694-A 79 12-DEC-2000;
FEATURES
source location/Qualifiers
1..20
BASE COUNT 9 a 3 c 3 g 5 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 173 attagataaaag 185
|||||
Db 2 ATTGATGATGAAAG 14

RESULT 24
LOCUS AX076045/c 20 bp DNA
DEFINITION Sequence 21 from Patent WO0104358.
ACCESSION AX076045
VERSION AX076045.1 GI:12710698
KEYWORDS
SOURCE Hepatitis B virus.
ORGANISM Hepatitis B virus.
REFERENCE 1 (bases 1 to 20)
AUTHORS Stuyver,L., Maertens,G. and van Geyt,C.
TITLE Detection of anti-hepatitis b drug resistance
JOURNAL Patent: WO 0104358-A 21 18-JAN-2001;
FEATURES
source location/Qualifiers
1..20
/organism="Hepatitis B virus"
/db_xref="taxon:10407"

BASE COUNT 6 a 3 c 4 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 72 tatatagtctgaaa 84
|||||
Db 15 TATATAGCTGTAAG 3

RESULT 25
LOCUS A97479 23 bp DNA
DEFINITION Sequence 35 from Patent WO9916780.
ACCESSION A97479
VERSION A97479.1 GI:6780825

KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.

REFERENCE 1 (bases 1 to 23)
AUTHORS Gala,J. and Vannuffel,P.
TITLE GENETIC SEQUENCES, DIAGNOSTIC AND/OR QUANTIFICATION METHODS AND DEVICES FOR THE IDENTIFICATION OF STAPHYLOCOCCI STRAINS
JOURNAL Patent: WO 9916780-A 35 08-APR-1999;
FEATURES
source location/Qualifiers
1..23
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 11 a 2 c 3 g 7 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 23;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 58 aaagatatataa 70
|||||
Db 6 AAAGATATATATTA 18

RESULT 26
LOCUS AX093544 24 bp DNA
DEFINITION Sequence 74 from Patent WO0118198.
ACCESSION AX093544
VERSION AX093544.1 GI:13509982
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 24)
AUTHORS Weissbach,J. and Hazan,J.
TITLE Cloning, expression and characterisation of the spg4 gene responsible for the most frequent form of autosomal spastic paraplegia
JOURNAL Patent: WO 0118198-A 74 15-MAR-2001;
FEATURES
source location/Qualifiers
1..24
/organism="Homo sapiens"
/db_xref="taxon:9606"
/note="Site accepteur d' passage du g ne spg4."

BASE COUNT 8 a 2 c 4 g 10 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 199 ttcttatlttaaa 211
|||||
Db 3 TTTTATATTTTAA 15

RESULT 27
LOCUS AX164353 24 bp DNA
DEFINITION Sequence 183 from Patent WO0138564.
ACCESSION AX164353
VERSION AX164353.1 GI:14545287
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequence.

REFERENCE 1 (bases 1 to 24)
 AUTHORS Rouleau,G.A., Lafreniere,R.G., Rochefort,D., Cossette,P. and Ragsdale,D.
 TITLE Loci for idiopathic generalized epilepsy, mutations thereof and method using same to assess, diagnose, prognosis or treat epilepsy
 JOURNAL Patent: WO 0138564-A 183 31-MAY-2001;
 McGill University (CA)
 FEATURES
 SOURCE location/Qualifiers
 1..24
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="synthetic oligonucleotide"
 BASE COUNT 2 a 5 c 3 g 14 t
 ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 301 ttatgtttcat 313
 |||||||||
 Db 8 TTTATGTTTCAT 20

RESULT 28
 AB069100/c 24 bp DNA SYN 08-AUG-2001
 LOCUS Synthetic construct DNA, forward primer for human STS-R133B1R
 DEFINITION at 1p36.
 ACCESSION AB069100
 VERSION AB069100.1 GI:15129904
 KEYWORDS
 SOURCE synthetic construct DNA.
 ORGANISM
 REFERENCE 1 (bases 1 to 24)
 AUTHORS Chen,Y.Z., Hayashi,Y., Wu,J.G., Takaoka,E., Maekawa,K., Watanabe,N., Inazawa,J., Hosoda,F., Arai,Y., Mizushima,H., Morohashi,A., Ohira,M., Nakagawara,A., Liu,S., Hoshi,M., Horii,A. and Soeda,E.
 TITLE A bac-based sts-content map spanning a 35-mb region of human chromosome 1p35-p36
 JOURNAL Genomics. 74 (1), 55-70 (2001)
 MEDLINE 21269192
 REFERENCE 2 (bases 1 to 24)
 AUTHORS Horii,A.
 TITLE Direct Submission
 JOURNAL Submitted (04-AUG-2001) Akira Horii, Tohoku University School of Medicine, Molecular Pathology; 2-1 Seiryomachi, Aoba-ku, Sendai, Miyagi 980-8575, Japan (E-mail:horii@mail.cc.tohoku.ac.jp, Tel:81-22-717-8042, Fax:81-22-717-8047)

FEATURES
 SOURCE location/Qualifiers
 1..24
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 misc_feature 1..24
 /note="forward primer for human STS-R133B1R at 1p36 sts-R133B1R obtained from clones B367G16, B133B1, B247E16, B132L12, B32A16, B76C16, B81K9, B32I16, Human BAC library RPCI-11"

BASE COUNT 9 a 2 c 4 g 9 t
 ORIGIN

Query Match 4.0%; Score 13; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 227 atataatgaatt 239
 |||||||||
 Db 14 ATATATGTAAT 2

RESULT 29
 AX042574/c 25 bp DNA PAT 23-NOV-2000
 LOCUS Sequence 140 from Patent WO0065088.
 DEFINITION AX042574
 ACCESSION AX042574
 VERSION AX042574.1 GI:11341182
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM
 REFERENCE 1 (bases 1 to 25)
 AUTHORS Ulfendahl,P.J. and Wong,K.C.
 TITLE Primers for identifying typing or classifying nucleic acids
 JOURNAL Patent: WO 0065088-A 140 02-NOV-2000;
 Amersham Pharmacia Biotech AB (SE)
 FEATURES
 SOURCE location/Qualifiers
 1..25
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="DQAI Homozygote primer sequence"
 BASE COUNT 4 a 5 c 3 g 13 t
 ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 237 attgtaaaaaa 249
 |||||||||
 Db 17 ATTGTAAAAAAA 5

RESULT 30
 AX043268/c 25 bp DNA PAT 23-NOV-2000
 LOCUS Sequence 834 from Patent WO0065088.
 DEFINITION AX043268
 ACCESSION AX043268
 VERSION AX043268.1 GI:11341876
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM
 REFERENCE 1 (bases 1 to 25)
 AUTHORS Ulfendahl,P.J. and Wong,K.C.
 TITLE Primers for identifying typing or classifying nucleic acids
 JOURNAL Patent: WO 0065088-A 834 02-NOV-2000;
 Amersham Pharmacia Biotech AB (SE)
 FEATURES
 SOURCE location/Qualifiers
 1..25
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="DQAI Heterozygote Primer Sequence"
 BASE COUNT 4 a 5 c 3 g 13 t
 ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 237 attgtaaaaaa 249
 |||||||||
 Db 17 ATTGTAAAAAAA 5

RESULT 31
 AX039624 26 bp DNA PAT 18-NOV-2000
 LOCUS Sequence 13 from Patent WO0063441.
 DEFINITION AX039624
 ACCESSION AX039624.1 GI:11229653
 VERSION

KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herrnstadt, C. and Davis, R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with Alzheimer's disease
JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
MITOKOR (US)

FEATURES
SOURCE Location/Qualifiers
1..26
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"

BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 304 atgtttcatgtt 316
|||||
Db 7 ATGTTTCATGTT 19

RESULT 32
AX039654 26 bp DNA PAT 18-NOV-2000
LOCUS AX039654
DEFINITION Sequence 43 from Patent WO0063441.
ACCESSION AX039654
VERSION AX039654.1 GI:11229683
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herrnstadt, C. and Davis, R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with Alzheimer's disease
JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
MITOKOR (US)

FEATURES
SOURCE Location/Qualifiers
1..26
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"

BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 304 atgtttcatgtt 316
|||||
Db 7 ATGTTTCATGTT 19

RESULT 33
AR014028 28 bp DNA PAT 05-DEC-1998
LOCUS AR014028
DEFINITION Sequence 64 from patent US 5773246.
ACCESSION AR014028
VERSION AR014028.1 GI:3971482
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 28)
/db_xref="taxon:32630"
/note="Artificial"

AUTHORS Keene, J.D., Levine, T. and Gao, F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5773246-A 64 30-JUN-1998;
FEATURES
SOURCE Location/Qualifiers
1..28
/organism="unknown"

BASE COUNT 6 a 3 c 1 g 18 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208
|||||
Db 3 TTATTTTATTTT 15

RESULT 34
I21978 28 bp DNA PAT 07-OCT-1996
LOCUS I21978
DEFINITION Sequence 64 from patent US 5525495.
ACCESSION I21978
VERSION I21978.1 GI:1602332
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 28)
AUTHORS Keene, J.D., Levine, T. and Gao, F.
TITLE Methods and compositions useful in the recognition, binding and expression of ribonucleic acids involved in cell growth, neoplasia and immunoregulation
JOURNAL Patent: US 5525495-A 64 11-JUN-1996;
FEATURES
SOURCE Location/Qualifiers
1..28
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BASE COUNT 6 a 3 c 1 g 18 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208
|||||
Db 3 TTATTTTATTTT 15

RESULT 35
AX012366 29 bp DNA PAT 06-SEP-2000
LOCUS AX012366
DEFINITION Sequence 26 from Patent EP0955369.
ACCESSION AX012366
VERSION AX012366.1 GI:9998412
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 29)
AUTHORS Bartscher, H.D., Mueller, R.D., Hoelke, W.D. and Millan, J.L.
TITLE High active alkaline phosphatase
JOURNAL Patent: EP 0955369-A 26 10-NOV-1999;
ROCHE DIAGNOSTICS GMBH (DE)

FEATURES
SOURCE Location/Qualifiers
1..29
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Artificial"

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BASE COUNT      8 a      7 c      9 g      5 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 29;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      22 gcacagaagaatg 34
        |||||||
        9 GCACAGAGAGATG 21

RESULT 36
E59972      29 bp      DNA      PAT      07-FEB-2001
LOCUS      Highly active alkaline phosphatase.
DEFINITION E59972
ACCESSION E59972.1 GI:13017742
VERSION JP 1999332586-A/23.
KEYWORDS JP 1999332586-A/23.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE unclassified.
          1 (bases 1 to 29)
AUTHORS Werner, H.R.M.M. and Burutosh, J.L.M.M.
TITLE Highly active alkaline phosphatase
JOURNAL Patent: JP 1999332586-A 23 07-DEC-1999;
          ROCHE DIAGNOSTICS GMBH
COMMENT OS Artificial Sequence
          PN JP 1999332586-A/23
          PD 07-DEC-1999
          PE 06-MAY-1999 JP 1999126494
          PR 05-MAY-1998 DE 19819962;7
          PI WERNER HOERKU, REINA MULLER, HERMUTTO BURUTOSHA, PI JOSE
          LOUIS MILAN
          PC C12N15/09, C12N1/15, C12N1/19, C12N1/21, C12N5/10, C12N9/16, PC
          C12N15/00, C12N5/00
          CC
          FH Key Location/Qualifiers
          FT source 1..29
          FT /organism='Artificial Sequence'
          FT Location/Qualifiers
          1..29
          /organism='unidentified'
          /db_xref='taxon:32644'

BASE COUNT      8 a      7 c      9 g      5 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 29;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      22 gcacagaagaatg 34
        |||||||
        9 GCACAGAGAGATG 21

RESULT 37
AR028230      30 bp      DNA      PAT      29-SEP-1999
LOCUS      Sequence 79 from patent US 5858661.
DEFINITION AR028230
ACCESSION AR028230
VERSION AR028230.1 GI:5940203
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE unclassified.
          1 (bases 1 to 30)
AUTHORS Shiloh, Y.
TITLE Ataxia-telangiectasia gene and its genomic organization
JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
          Location/Qualifiers

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source 1..30
/organism="unknown"
BASE COUNT      13 a      2 c      1 g      14 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      272 aaaaattatc 284
        |||||||
        15 AAAAATTATTC 27

RESULT 38
AR138633      30 bp      DNA      PAT      16-JUN-2001
LOCUS      AR138633
DEFINITION Sequence 158 from patent US 6200749.
ACCESSION AR138633
VERSION AR138633.1 GI:14480978
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE unclassified.
          1 (bases 1 to 30)
AUTHORS Shiloh, Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to
          screen for a partial A-T phenotype
JOURNAL Patent: US 6200749-A 158 13-MAR-2001;
          Location/Qualifiers
          1..30
          /organism="unknown"

BASE COUNT      13 a      2 c      1 g      14 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      272 aaaaattatc 284
        |||||||
        15 AAAAATTATTC 27

RESULT 39
AX063379      30 bp      DNA      PAT      24-JAN-2001
LOCUS      AX063379/C
DEFINITION Sequence 42 from Patent WO0079009.
ACCESSION AX063379
VERSION AX063379.1 GI:12541169
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
          artificial sequence.
REFERENCE 1 (bases 1 to 30)
AUTHORS Nazarenko, I. and Rashtchian, A.
TITLE Improved primers and methods for the detection and discrimination
          of nucleic acids
JOURNAL Patent: WO 0079009-A 42 28-DEC-2000;
          Life Technologies, Inc. (US)
          Location/Qualifiers
          1..30
          /organism="synthetic construct"
          /db_xref='taxon:32630'
          /note="Oligonucleotide"

BASE COUNT      10 a      4 c      7 g      9 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 13; DB 6; Length 30;

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Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 40 atgaataaagat 52
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Db 14 ATGAATAAAGAT 2

RESULT 40
LOCUS 195122/c 31 bp DNA PAT 01-DEC-1998
DEFINITION Sequence 5 from patent US 5731425.
ACCESSION 195122
VERSION 195122.1 GI:3939592
KEYWORDS
SOURCE .
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 31)
Brizzard,B.L., Blanca,D.W., Chubet,R.G., Vizard,D.L. and
Hopp,T.Patrick.
TITLE Polypeptide surface marker for cells
JOURNAL Patent: US 5731425-A 5 24-MAR-1998;
FEATURES Location/Qualifiers
source 1..31
BASE COUNT 12 a 11 c 7 g 1 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 31;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 124 gtgtctctcgagg 136
|||||
Db 28 GTGTTCTCGGGG 16

RESULT 41
XELARSE59 33 bp DNA VRT 28-APR-1993
LOCUS XENOPUS laevis autonomous replication sequence e59.
DEFINITION R01606
ACCESSION R01606.1 GI:213953
VERSION
KEYWORDS autonomous replication; mutational analysis.
SOURCE XENOPUS laevis DNA.
ORGANISM XENOPUS laevis
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipridae; Pipidae;
Xenopodinae; Xenopus.

REFERENCE 1 (bases 1 to 33)
AUTHORS Kearsey,S.
TITLE Structural requirements for the function of a yeast chromosomal
JOURNAL replicator
MEDLINE Cell 37, 299-307 (1984)
FEATURES 84205653
source Location/Qualifiers
1..33
/organism="Xenopus laevis"
/db_xref="taxon:8355"

BASE COUNT 8 a 4 c 2 g 19 t
ORIGIN

Query Match 4.0%; Score 13; DB 5; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 298 attttatgttt 310
|||||
Db 1 ATTATTATGATTT 13

RESULT 42
LOCUS A41027/c 36 bp DNA PAT 05-MAR-1997
DEFINITION Sequence 14 from Patent EP0626449.
ACCESSION A41027
VERSION A41027.1 GI:2296916
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified

REFERENCE 1 (bases 1 to 36)
AUTHORS Schreier,P.H., Stenzel,K.D., Adam,G.P. and Maiss,E.D.
TITLE DNA coding for plant virus sequences
JOURNAL Patent: EP 0626449-A 14 30-NOV-1994;
BAYER AG (DE)
COMMENT Other publication JP 6343469 941220
Other publication CA 2124272 941129
Other publication CN 1098744 950215
Other publication AU 6191494 941201
Other publication DE 4317845 941201
Other publication ZA 9403730 950202.
FEATURES Location/Qualifiers
source 1..36
/organism="unidentified"
/db_xref="taxon:32644"

BASE COUNT 9 a 5 c 5 g 17 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 156 aggaataataaa 168
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Db 24 AGGAATAATATAA 12

RESULT 43
AR082586/c 36 bp DNA PAT 31-AUG-2000
LOCUS AR082586
DEFINITION Sequence 14 from patent US 5973135.
ACCESSION AR082586
VERSION AR082586.1 GI:10009308
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 36)
AUTHORS Schreier,P.Helmolt, Stenzel,K., Adam,G. and Maiss,E.
TITLE DNA comprising plum pox virus and tomato spotted wilt virus CDNAS
JOURNAL for disease resistance
MEDLINE Patent: US 5973135-A 14 26-OCT-1999;
FEATURES Location/Qualifiers
source 1..36
/organism="unknown"
BASE COUNT 9 a 5 c 5 g 17 t
ORIGIN

Query Match 4.0%; Score 13; DB 6; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 156 aggaataataaa 168
|||||
Db 24 AGGAATAATATAA 12

RESULT 44
AXI67671 36 bp DNA PAT 03-JUL-2001
LOCUS AXI67671
DEFINITION Sequence 16 from Patent WO0144277.

ACCESSION AX167671 GI:14597058
 VERSION AX167671.1

KEYWORDS
 SOURCE synthetic construct.
 ORGANISM synthetic construct.
 artificial sequence.

REFERENCE 1 (bases 1 to 36)

AUTHORS Weglich Glover, L., Budziszewski, G. J., Levin, J. Z., and Zhou, Q.

TITLE Herbicide target genes and methods

JOURNAL Patent: WO 0144277-A 16 21-JUN-2001;

FEATURES
 source Syngenta Participations AG (CH)
 Location/Qualifiers
 1..36

BASE COUNT /organism="synthetic construct"
 ORIGIN /db_xref="taxon:32630"
 6 a 4 c 11 g 15 t
 4 c 11 g 15 t

Query Match 4.0%; Score 13; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 303 tatgtttcatgt 315

DB 21 TATGTTTTCATGT 33

RESULT 45

LOCUS 128261 36 bp DNA PAT 06-FEB-1997

DEFINITION Sequence 14 from patent US 5569823.

ACCESSION 128261

VERSION 128261.1 GI:1819037

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 36)

AUTHORS Schreier, P. H., Stenzel, K., Adam, G., Unter and Maiss, E.

TITLE DNA comprising plum pox virus and tomato spotted wilt virus cDNAs

JOURNAL Patent: US 5569823-A 14 29-OCT-1996;

FEATURES Location/Qualifiers

source 1..36

BASE COUNT /organism="unknown"
 ORIGIN 9 a 5 c 5 g 17 t

Query Match 4.0%; Score 13; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 156 aggaataataaa 168

DB 24 AGGAATAATATAA 12

Search completed: January 24, 2002, 03:22:36
 Job time: 3813 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:12:27 ; Search time 1502.85 Seconds
(without alignments)

3589.565 Million cell updates/sec

Title: US-09-531-438-3

Perfect score: 327

Sequence: 1 attggagatacttaattt.....tttcattttctattgtt 327

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1472140 seqs, 8248589755 residues

Total number of hits satisfying chosen parameters: 2944280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

GenBml:*

1: gb_ba:*
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4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_om:*
20: em_or:*
21: em_ov:*
22: em_pat:*
23: em_ph:*
24: em_pl:*
25: em_ro:*
26: em_sts:*
27: em_sy:*
28: em_un:*
29: em_vi:*
30: em_htgo_hum:*
31: em_htgo_inv:*
32: em_htgo_rod:*
33: em_htg_hum:*
34: em_htg_inv:*
35: em_htg_rod:*
36: em_htg_other:*

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	327	100.0	327	6 AX004614	AX004614 Sequence
2	327	100.0	1392	1 L77965	L77965 Clostridium
3	327	100.0	1392	6 AX004613	AX004613 Sequence
4	199	50.9	54310	1 AP003515	AP003515 Clostridi
5	81.8	25.0	12900	3 AE001429	AE001429 Plasmodiu
6	79	24.2	39347	9 AL135906	AL135906 Human DNA
7	77.8	23.8	53932	2 AC023371	AC023371 Homo sapi
8	77.4	23.7	163443	2 AC006280	AC006280 Plasmodiu
9	77.4	23.7	205429	2 AC005506	AC005506 Plasmodiu
10	76.2	23.3	3392	2 AF300334	AF300334 Dictyoste
11	76.2	23.3	242513	2 AC079314	AC079314 Homo sapi
12	75.8	23.2	318221	2 PFMAL13P3	PFMAL13P3 Plasmodiu
13	75.6	23.1	140414	2 AF377947	AF377947 Oriza sat
14	75	22.9	178783	2 AC068139	AC068139 Homo sapi
15	74.4	22.8	156060	2 AC004153	AC004153 Plasmodiu
16	73.2	22.4	863	11 CNS06EVO	AL395628 T7 end of
17	73	22.3	8622	8 YSCMTCTOC	M97514 Saccharomyc
18	72.8	22.3	34119	8 AF222718	AF222718 Chrysoid
19	72.2	22.1	104992	2 AC005504	AC005504 Plasmodiu
20	72.2	22.1	162445	2 AL158151	AL158151 Human DNA
21	72.2	22.1	169546	2 AC004157	AC004157 Plasmodiu
22	72.2	22.1	199882	2 AL354720	AL354720 Human DNA
23	72	22.0	158398	2 AC011146	AC011146 Homo sapi
24	71.6	21.9	180903	3 AC073409	AC073409 Homo sapi
25	71.4	21.8	12029	3 AE001400	AE001400 Plasmodiu
26	71.4	21.8	175053	2 AC090014	AC090014 Homo sapi
27	71.2	21.8	110000	2 AL591074-2	Continuation (3 of
28	71	21.7	95477	9 AC007076	AC007076 Homo sapi
29	71	21.7	168799	9 AC009531	AC009531 Homo sapi
30	71	21.7	194038	9 AC010103	AC010103 Homo sapi
31	70.8	21.7	159475	2 AC021378	AC021378 Homo sapi
32	70.6	21.6	13433	3 AF315648	AF315648 Ceratilis
33	70.6	21.6	85779	8 SCE011856	AU011856 Saccharom
34	70.6	21.6	122747	2 AC093220	AC093220 Homo sapi
35	70.6	21.6	159255	9 AF212831	AF212831 Homo sapi
36	70.6	21.6	161230	2 AC011355	AC011355 Homo sapi
37	70.6	21.6	234112	3 PFMAL4P2	AL035475 Plasmodiu
38	70.6	21.6	340000	2 HS21C013	AL163213 Homo sapi
39	70.4	21.5	155456	2 AC027753	AC027753 Homo sapi
40	70.4	21.5	160624	9 AC060835	AC060835 Homo sapi
41	70.4	21.5	172758	2 AC022553	AC022553 Homo sapi
42	70.4	21.5	195551	2 AC006281	AC006281 Plasmodiu
43	70.2	21.5	137342	9 AL392048	AL392048 Human DNA
44	70.2	21.5	180388	9 HUMRETLAS	L11910 Human retin
45	70.2	21.5	183584	9 AC012492	AC012492 Homo sapi

ALIGNMENTS

RESULT 1
AX004614 LOCUS AX004614 327 bp DNA
DEFINITION Sequence 2 from Patent WO9915669.
ACCESSION AX004614
VERSION AX004614.1 GI:9928055
KEYWORDS
SOURCE
ORGANISM

Clostridium perfringens.
Clostridium perfringens
Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;

REFERENCE 1 (bases 1 to 327)

AUTHORS Gilbert,M. and Popoff,M.R.
TITLE Clostridium toxin, and method for preparing immunogenic
compositions

JOURNALS Patient: WO 9915669-A 2 01-APR-1999;
GIBERT MARISE (FR); PASTEUR INSTITUT (FR)
FEATURES
source
1. 327
Location/Qualifiers

/organism="Clostridium perfringens"

[/db_xref="GI:9328054"](#)
[/translation="MKIISKFTVIFMFSCLLIVGAIISPMKASKEIDAYRKMENTE](#)

NALNKNDINTVNISEDERNVNNVQYREMLEDEKYPDNQOLKSEFILLNSOKDNKEIF
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 AQAIGFYQPTQYDREFTYVADNILLNFRQVATSGSRDLKVEYSVVDHMKKDKVKASQ
 MVGQNPDSKRQRLRIETKQCSYKRIKRIKFTPASIRVFGEJCA

BASE COUNT 606 a 115 c 209 g 462 t
 ORIGIN

Query Match 100.0%; Score 327; DB 6; Length 1392;
 Best Local Similarity 100.0%; Pred. No. 2.8e-28;
 Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 attgggagatccttaaatagcacaagaagtgttaaatgaaataaagaataaataa 60
 Db 1 ATTTGGGATCTTAAATTAGCACAGAAATGTTAAATGAATTAAGTAATAA 60
 Qy 61 gatataatataatagctgaataattataatataatgaatagtaataataa 120
 Db 61 GATATATTAATTAATAGCTGAAATTTATATATATAGTAATAGTAATAATA 120
 Qy 121 aaagtgcttcgggggacactttttgtttaaaaaggaataataataattagat 180
 Db 121 AAAGTGCTTCGGGGGACACTTTTGTTTTAAAGGAAATATATAAATTAATGAT 180
 Qy 181 aaagtgaataataatttttatttaattgttaaaaattgataataattgaaatg 240
 Db 181 AAAAGTGAATAATTAATTTTATTTTAAATTGTTAAATTTGATATATGAAATG 240
 Qy 241 taaaaaaattcaggggggaataataatgaataaataattcagaattactgtaatt 300
 Db 241 TAAAAAAATTTACGGGGGGAATTAATGAATAAATTAATTTCAAGTTACTGTAAT 300
 Qy 301 ttatgttttcattgtttttctatcgtt 327
 Db 301 TTATGTTTTCATGTTTTCATATGTT 327

RESULT 4
 LOCUS AP003515 54310 bp DNA circular BCT 10-AUG-2001
 DEFINITION Clostridium perfringens plasmid pCP13 DNA, complete sequence.
 ACCESSION AP003515
 VERSION AP003515.1 GI:15076712
 KEYWORDS
 SOURCE Clostridium perfringens (strain:13) plasmid:pCP13 DNA.
 ORGANISM Clostridium perfringens
 Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;

REFERENCE 1 (bases 1 to 54310)
 AUTHORS Ohtani,K., Ohshima,S., Hirakawa,H., Ohshima,K., Shiba,T.,
 Shimizu,T., Hattori,M., Kuhara,S., Hayashi,H. and Shimizu,T.
 TITLE Complete Nucleotide Sequence of the Virulence Plasmid pCP13 from
 Clostridium perfringens
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 54310)
 AUTHORS Shimizu,T.
 JOURNAL Direct Submission
 TITLE Submitted (12-APR-2001) Tohru Shimizu, Institute of Basic Medical
 Sciences, University of Tsukuba, Department of Microbiology; 1-1-1
 Tennoudai, Tsukuba, Ibaraki 305-8575, Japan
 E-mail:tschimizuend.tsukuba.ac.jp, Tel:81-298-53-3354,
 Fax:81-298-53-3354)

FEATURES
 source
 1. 54310
 location/Qualifiers
 /organism="Clostridium perfringens"
 /plasmid="pCP13"
 /strain="13"
 /db_xref="taxon:1502"
 /note="anaerobic pathogen for gas gangrene"
 940..1692
 /gene="soj"
 940..1692
 /gene="soj"
 /gene="soj"

gene
 CDS

/note="250 aa, similar to p1r:14044 Sp00A activation
 inhibitor soj from Bacillus subtilis (253 aa); 378
 identity in 250 aa overlap
 pCP01

Para family"
 /codon_start=1
 /transl_table=11
 /product="Soj protein"
 /protein_id="BAB62438.1"
 /db_xref="GI:15076713"
 /translation="MKISVFNKGGVAKTTPANGACLEKGRVLLVLDPOSNL
 TKLFKAVSMEDVSIADVLIDKNDIEKVIKTPFENDIPSVTLAFARKILLDVN
 RSQONRLAKLEIEDKDYDCLIDCPALMTITVNLACASDEVLPKIDPKLDGLE
 YLDSIEERKDEFNPINFEKCFITMDSSTTVKVKYKQELKSVLGEKMRNTSIHQNIK
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 1751..3031
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 1751..3031
 /gene="parB"
 /note="426 aa, similar to gpu:AF300944_3 presumptive ParB
 protein from Lactococcus lactis subsp.lactis (242 aa); 308
 identity in 266 aa overlap
 pCP02"

/codon_start=1
 /transl_table=11
 /product="ParB protein"
 /protein_id="BAB62439.1"
 /db_xref="GI:15076714"
 /translation="MAKFSISSEGMNGISKNTRKRYEEQAKENFTEVINDIRKN
 EKNFEIVDIESLAEDIKINGLNHNLVVRKLDMDVELISGERRYALSKLVEGNKE
 FNLVPCKVIESNDISEIILIOANQOSRELTEVEKLTVERLOEYFKIRKENGKVP
 KIRDIIDLNLSTATQVGRERYNNKRLIPEIKAVITBOGNTTANASEPSSISENRY
 ILSIIDKTRMSQEAVDLKNKRLKIEBEKELELKKAYEKELELRLBEKKNQVSK
 LKSENLKRLKDSNNIIEBKETBGQIKIEFEKLENEKVIIEELKSKYDKKIEDI
 TEAKENNLKORLKDLELSLKRSENEVDIKTKENFVLVONLKLIDNSFKMLKSOI
 NKKKENVVVAEEETKAKEPLEKYKEISDLKTL"
 3147..3509
 /gene="pCP03"
 3147..3509
 /gene="pCP03"
 /note="120 aa, no significant homology"
 /codon_start=1
 /transl_table=11
 /product="hypothetical protein"
 /protein_id="BAB62440.1"
 /db_xref="GI:15076715"
 /translation="MEKILAERKINISFYKRRKGAALVTLIYPPKMLEVIGITENERE
 CFEYIDKAIKISKERQSEAKERTISFSTSTKTYLNNKMLEYLGVSDEDSCTIEL
 RKKIDILVRKNGRDIDDI"
 3773..4024
 /gene="pCP04"
 3773..4024
 /gene="pCP04"
 /note="83 aa, similar to p1r:T14710 probable transposase
 from Yersinia pestis (402 aa); 44% identity in 50 aa
 overlap
 truncated"

/codon_start=1
 /transl_table=11
 /product="probable transposase"
 /protein_id="BAB62442.1"
 /db_xref="GI:15076717"
 /translation="MSKDFLRNYIKRQNFNSNNVLSIKDLFNDVLOEVSAELDDM
 LGYEKLITNOFYRNRGYSKRTIKSELGTTITLILRYRN"
 4040..4422
 /gene="pCP05"
 4040..4422
 /gene="pCP05"
 /note="60 aa, similar to gp:AF143819.1 transposase-like
 protein from Escherichia coli (402 aa); 38% identity in 60
 aa overlap
 truncated"
 /codon_start=1

gene
CDS
/transl_table=1
/product="probable transposase"
/protein_id="BAB62443.1"
/db_xref="GI:15076718"
/translation="MISKHQRNINRIEDKNLNLVASCITTRDVAGQIKALYDIEISAE
TVSNITNIMPLVSEW"
4588..4746
/gene="PCP06"
4588..4746
/note="52 aa, similar to p1r:T43600 probable transposase
from Yersinia pestis (105 aa); 56% identity in 50 aa
overlap
truncated"
/codon_start=1
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/product="probable transposase"
/protein_id="BAB62444.1"
/db_xref="GI:15076719"
/translation="MLYTTNVIESLNSQFRKFTKTLIFPNDVSLIKMLYLATEKVNK
KWRPNYPR"
complement(5169..5804)
/gene="PCP07"
complement(5169..5804)
/gene="PCP07"
/function="ATP-binding protein"
/note="211 aa, similar to gp:AB001508.4 ABC transporter
(ATP-binding protein) from Bacillus halodurans (213 aa);
49% identity in 214 aa overlap"
/codon_start=1
/transl_table=1
/product="probable ABC transporter"
/protein_id="BAB62445.1"
/db_xref="GI:15076720"
/translation="MNTIEISNLKKRYFDKIFKDFSLISIKGEMIAISGSGCGKST
LNLMLGLEKFDSEIITDGVKNIKNSLANKFLREKISTYLFQNFALVDELYEENL
RLAKHTIKNTKRIEELIRCLKFGLEGCKKNYIELSGEQORVALARMLKPEEL
ILADEPTGSLDENRDIISLKLKELNESGKTIITVHDNVAKADRITFL"
complement(5804..7966)
/gene="PCP08"
complement(5804..7966)
/note="720 aa, similar to gp:AB001508.3 BH0280 gene
product from Bacillus halodurans (713 aa); 23% identity in
661 aa overlap"
/codon_start=1
/transl_table=1
/product="conserved hypothetical protein"
/protein_id="BAB62446.1"
/db_xref="GI:15076721"
/translation="MKKKVALIILFIITLISFGVYSVRNHTEFMKLNLOCNENF
EVTISIDREHNATVASEIISKSLDEYNITFSEVDIEDVAKYINSGFENKENEH
IPVSGRFHEKNDISYLTIDSEDTQICVINDFNKELHEITLTKSDINTFEN
LFIQIENEQILDKLIEDLKSESIIVQKVGSDSSQYETELKILLVNCFGLFMIF
YQVLSGYKIKIGIKLGHSTFVLMLEKLELVLEIVLMLVTVTLIIVNCFGLFMIF
WKFMELICISIMIFIVISVPIYIVSKITSLINKRPVKSIIILNKSIVAIL
ASLIFPSNALDLSISIGKEKNYKWEETKOYILPELGPNDSESTOSPIEMEK
ERAYIYVNGKATILADPNRYEPREMEAKOMLPEYVRETIIVNPNLKHKYVD
GNITISDEKDRILVPEKTRNEKELELYGINSQPEGSSTTCSHTAGCKMLVE
QKQKTIWMSNQKFEYSLDVNPEGNVTPPIVSVLESNDKLVSYKIIQYNNSP
KIRNSEEVEVNGLEKYDMGVYLDIPNLDVNASIINKAKKVIIFIVILLAV
ISIILOMTSLYFNQNNKRIYVKLHGVRILRYKMYPIFIMVLITWTCPLAASLITKD
INIITYLIVVIEVFIFENINSLEKKNLKVIGKEV"
complement(8043..8354)
/gene="PCP09"
complement(8043..8354)
/note="103 aa, no significant homology"
/codon_start=1
/transl_table=1
/product="hypothetical protein"
/protein_id="BAB62447.1"
/db_xref="GI:15076722"

gene
CDS
/translation="MKRNLIKTFIASLLVGSIGTATALAYTHSSDKFEASLPLGFS
QAQSSKFFYCGGQKRIKATATARYKVGTTLVEANKIKAKILAHNQCTYYVGVEMNSYVYAH
L"
8779..9012
/gene="PCP10"
8779..9012
/note="77 aa, similar to probable transposase from
Yersinia pestis plasmid pMT1 (402 aa); 25% identity in 158
aa overlap
truncated"
/codon_start=1
/transl_table=1
/product="probable transposase"
/protein_id="BAB62448.1"
/db_xref="GI:15076723"
/translation="MKEVLSTCMSTNEGSKFWLSPHKDKKEFAKDLKTIYGSVNETGS
MKNDLEIREKWSKYPNVVKSWMKDNMDNLSTFF"
9127..9366
/gene="PCP11"
gene
Query Match 60.9%; Score 199; DB 1; Length 54310;
Best Local Similarity 86.3%; Pred. No. 2.2e-14;
Matches 220; Conservative 0; Mismatches 35; Indels 0; Gaps 0;
QY 73 atatagctgaataattatataatgataagatgaatgaataaagaagtgcttcg 132
DB 13459 ATGAAATTTTAACTCATGCTTTTAAAGTTAATATATATTTTAAATTAGAGTGCCAC 13518
QY 133 gggagacacttttgctttaaaaggaaataataaattagataaaggatgataa 192
DB 13519 GGGGACACTCTTTTGCTTTTAAAGTAAATATGATTAATTTAGATTAAGTGTAA 13578
QY 139 taattatttattttaaattgttaaaattgataaattgataaattgtaaaaaaatt 252
DB 13579 GAATTAATTTTAAATTTTAAATTTGTTAAATTTGATTAATGATTTGTAATAAATTT 13638
QY 253 cagggggggaataataaataaataattcaaaattcaagttactgtaattttagtttca 312
DB 13639 CAGGGGGGGAATTAATTAATAAATTTATTTCAAGTTTACTGTAATTTTATGTTTCA 13698
QY 313 tgtttcttattgtt 327
DB 13699 TATTTTCTTATGTGT 13713
RESULT 5
AE001429
LOCUS AE001429 12900 bp DNA INV 06-NOV-1998
DEFINITION Plasmodium falciparum chromosome 2, section 66 of 73 of the
complete sequence.
ACCESSION AE001429 AE001362
VERSION AE001429.1 GI:3845321
KEYWORDS
SOURCE
ORGANISM
Plasmodium falciparum.
REFERENCE
1 (bases 1 to 12900)
Gardner,M.J., Tetteilin,H., Carucci,D.J., Cummings,L.M., Aravind,L.,
Koonin,E.V., Shalim,S., Mason,T., Yu,K., Fujii,C., Pederson,J.,
Shen,K., Jing,J., Aston,C., Lai,D., Schwartz,D.C., Pertea,M.,
Salzberg,S., Zhou,L., Sutton,G.G., Clayton,R., White,O.,
Smith,H.O., Fraser,C.M., Hoffman,S.L., et.al.
Chromosome 2 sequence of the human malaria parasite Plasmodium
falciparum
JOURNAL Science 282 (5391), 1126-1132 (1998)
MEDLINE 99021743
REMARK Erratum: [[published erratum appears in Science 1998 Dec
4;282(5395):1827]]
2 (bases 1 to 12900)
Gardner,M.J.
AUTHORS Direct Submission
TITLE Submitted (02-NOV-1998) The Institute for Genomic Research, 9712

Medical Center Drive, Rockville, MD 20814, USA

FEATURES
Location/Qualifiers
1..12900

source
/organism="Plasmodium falciparum"
/db_xref="taxon:5833"
/chromosome="2"
11045..12083

gene
/gene="PFB0955w"
join(11045..11113,11229..12083)

CDS
/gene="PFB0955w"
/note="Identified by sequence similarity; putative"

/codon_start=1
/product="rflin"
/protein_id="AAC71980.1"
/db_xref="GI:3845322"

/translation="MNTYIMLMVLSLLVLEISYVNNHNKYNNGYIONNFQIM
KSRRLAEIQLPKCPHYNDPELKIIDKLEERIKRYIFENNSFEELHGLVERK
SLYENGKSSNMEKELIKKYDSDIRDEHNVISKGIYSRVLRYSCDYONKIL
RDELASCCYHDNDLNLKKGCFGVGICITCLSLVNSIGVYTAKEVITGLYSL
DIANKFTKLAGIYFFPSSISENGAGSVITFYWDSMRMASIASSTINPYGIALVLI
VLVVLYLVYIMVLRKKKSMKHECKHLS"

BASE COUNT 5356 a 845 c 1167 g 5532 t
ORIGIN

Query Match 25.0%; Score 81.8; DB 3; Length 12900;
Best Local Similarity 54.0%; Pred. No. 0.25; Mismatches 142; Indels 0; Gaps 0;

Matches 167; Conservative 0; Mismatches 142; Indels 0; Gaps 0;

OY 2 ttgggatacttaattagacagaagaatgttaaaatgaataaagaataaag 61

DB 6596 TTTCCGTTTCCTATATATATATACAGCGGTTAATTTATTTATTAATGAAT 6655

OY 62 atatttaattataagctgaatattataatataagctagctataataa 121

DB 6656 CTAATTTTATTTATTTATTAATAATATACATATATTTAATGCTAAACATATTA 6715

OY 122 aagtgctcggggagacctttgttttaaaaggaataataaataatagata 181

DB 6716 TAAATATATTTATATATATTTTATTTATTTATTAATAATGCAATATCATTTTGA 6775

OY 182 aaagtgaataaataattattatttaattgttaaaatgataataatgaatgt 241

DB 6776 ATAAAGAAATTTTATTTTATTTTATTTTAAATAATATATTTTAAACCTT 6835

OY 242 aaaaaaatttcagggggaataataaagaaataattcaaatctcgttaatt 301

DB 6836 TATAAATTAATAAATTTCTAATTAATAAATAATTTTAAACAATTTTATTAAT 6895

OY 302 ttatgttt 310

DB 6896 GAATATGTT 6904

RESULT 6
AL135906 39347 bp DNA PRI 26-APR-2000

LOCUS Human DNA sequence from clone RPI-20N11 on chromosome 6 Contains

ACCESSION AL135906
VERSION AL135906.19 GI:7248200

KEYWORDS
SOURCE human

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

AUTHORS
JOURNAL

Submitted (14-APR-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
Requests: clonerequest@sanger.ac.uk

COMMENT On Mar 15, 2000 this sequence version replaced gi:7210154.

FEATURES

source

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr6> RPI-20N11 is from the library RPI-1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/> VECTOR: pcypac2

IMPORTANT: This sequence is not the entire insert of clone RPI-20N11. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true right end of clone RPI-20N11 is at 39347 in this sequence. The true right end of clone RPI-482U3 is at 103 in this sequence.

Location/Qualifiers

1..39347

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="6"

/clone="RPI-20N11"

/clone_1lb="RPI-1"

2..1136

/note="L1P12 repeat: matches -1421..-291 of consensus"

3460..3893

/note="L1M3C repeat: matches 999..1402 of consensus"

3894..4213

/note="AluY repeat: matches 1..311 of consensus"

4214..4415

/note="L1M3C repeat: matches 1402..1610 of consensus"

4426..4777

/note="MER47A repeat: matches 13..366 of consensus"

4779..5831

/note="L1M2 repeat: matches 1528..2663 of consensus"

6028..6073

/note="L1 repeat: matches 2729..2777 of consensus"

6091..8167

/note="L1P15 repeat: matches 4085..6157 of consensus"

8287..8567

/note="AluX repeat: matches 3..284 of consensus"

8792..8965

/note="L1 repeat: matches 2737..2901 of consensus"

8959..9103

/note="L1M3C repeat: matches 1092..1236 of consensus"

9227..10104

/note="L1 repeat: matches 2903..3922 of consensus"

10091..10657

/note="L1M2 repeat: matches 5115..5687 of consensus"

10657..11436

/note="L1P12 repeat: matches 3..776 of consensus"

11432..16672

/note="L1P12 repeat: matches 900..6146 of consensus"

16682..16795

/note="L1P3 repeat: matches 5677..5791 of consensus"

16821..16860

/note="20 copies 2 mer ta 90% conserved"

16865..17035

/note="MER42 repeat: matches 2..152 of consensus"

```

repeat_region 17036..17343 /note="MER4A repeat: matches 367..646 of consensus"
repeat_region 17377..17678 /note="LIMA9 repeat: matches 5783..6112 of consensus"
repeat_region 17679..17996 /note="AluYb8 repeat: matches 1..318 of consensus"
repeat_region 17997..18193 /note="LIMA9 repeat: matches 6112..6300 of consensus"
repeat_region 19432..20219 /note="LIME repeat: matches 429..1193 of consensus"
misc_feature 23091..23131 /note="match: GSS: Em:AQ218253"
repeat_region 23144..23291 /note="MLTII repeat: matches 370..409 of consensus"
repeat_region 25530..25553 /note="LIPa4 repeat: matches 5999..6146 of consensus"
repeat_region 25570..25754 /note="12 copies 2 mer ca 95% conserved"
repeat_region 25827..26622 /note="AluIo/RRM repeat: matches 173..291 of consensus"
misc_feature 26639 /note="398 copies 2 mer aa 58% conserved"
misc_feature 25879..26639 /note="Tandem repeat. Tandem repeat contains forced join. BamI and EcoRI bot suggest approx 200bp missing."
misc_feature 26730 /note="match: GSS: Em:B69019"
misc_feature 26664 /note="match: GSS: Em:AQ230058"
repeat_region 27019..27243 /note="LTRA4 repeat: matches 282..516 of consensus"
repeat_region 27306..28098 /note="LIME2 repeat: matches 5094..5922 of consensus"
repeat_region 28440..28705 /note="133 copies 2 mer aa 54% conserved"
repeat_region 29274..29834 /note="LIMB6 repeat: matches 5552..6128 of consensus"
repeat_region 29885..31038 /note="LIM4 repeat: matches 862..2044 of consensus"
repeat_region 31742..32332 /note="LIMD1 repeat: matches 5317..5893 of consensus"
misc_feature 32769..32872 /note="match: GSS: Em:AQ078965"
repeat_region 32769..32872 /note="52 copies 2 mer tt 64% conserved"
misc_feature 33057 /note="match: GSS: Em:AQ780759"
repeat_region 33046..33127 /note="41 copies 2 mer tt 73% conserved"
repeat_region 33143..34362 /note="LIPa7 repeat: matches 4855..6127 of consensus"
repeat_region 34364..34625 /note="31 copies 2 mer ta 71% conserved"
repeat_region 34637..34664 /note="14 copies 2 mer ac 89% conserved"
repeat_region 34745..36725 /note="LIPa13 repeat: matches 2266..4198 of consensus"
repeat_region 36726..37032 /note="AluY repeat: matches 3..308 of consensus"
repeat_region 37033..38914 /note="LIPa13 repeat: matches 4198..6145 of consensus"
repeat_region 39090..39341 /note="TIGER2 repeat: matches 2448..2717 of consensus"

```

Db	25934	ATAAATATATATAAATATATATATATATATAAAATATATATATAAAATATATATAAAAT	25993
Oy	84	aattataaataatgatgaagtaagttaataaacaagaagtgctccggggacattc	143
Db	25994	ATATATATATATATAAAATATATATAAAATATATATAAAATATATATATAAAATAT	26053
Oy	144	tttgcttcaaaaggaatatataataaatctagataaagtcgtaaatcaattc	203
Db	26054	ATATATAATATATATAAAATATATATATATATATAAAATATATATAAAATATATATAT	26113
Oy	204	attttaaatttgtaaaaaatttgataaatctgaattgttaaaaaaaattcaggggggaat	263
Db	26114	ATAAATATATATAAAATATATATATATATAAAATATATAAAATATATATAAAATAT	26173
Oy	264	ataaatgaaaaaatattctcaaggtctacgtaatttttgcttc	310
Db	26174	ATTATATATATAAAATATATATAAAATATATATAAAATATATATAT	26220

	RESULT	7	
AC023371 LOCUS	AC023371	53932 bp	DNA HTG 13-JUL-2000
DEFINITION	Homo sapiens clone RP11-21D18,	LOW-PASS SEQUENCE SAMPLING.	
ACCESSION	AC023371		
VERSION	AC023371.2	GI:9123990	
KEYWORDS	HTG; HTGS_PHASE0.		
SOURCE	human.		

REFERENCE
AUTHORS
Birren, B., Linton, L., Nusbaum, C. and Lande, E.
1 (bases 1 to 53932)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

TITLE	Homo sapiens, clone RP11-21D18
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 53932)
AUTHORS	Birren, B., Linton, J., Nusbaum, C., Lande, E., Abraham, H., Allen, N.,

Anderson, S., Baldwin, J., Barua, N., Beda, F., Bonushavily, L.,
Boukhalter, B., Brown, A., Burkett, G., Campoliano, A., Castle, A.,
Choepl, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P.,
Dearellano, K., Dewar, K., Dodge, S., Domino, M., Doyle, M.,
Penestor, J., Ferreira, P., Fitzhugh, W., Forrest, C., Gage, D.,
Galagan, J., Gardyna, S., Ginde's, Goyette, M., Graham, L.,
Grand-Pierre, N., Grant, G., Haagos, B., Heatford, A., Horton, L.,
Howland, J. C., Iliev, I., Johnson, R., Jones, C., Kamm, L., Karataas, A.,
Klein, J., Landers, T., Largoque, K., Lehoczy, J., Levine, R.,
Lieu, C., Liu, G., Locke, K., Macdonald, P., Margus, N., McCarthy, M.,
McEwen, P., McGurk, A., McKernan, K., McPheeters, R., Meldrum, J.,
Meneus, L., Mihova, T., Miranda, C., Mlenga, V., Morrow, J., Naylor, J.,
Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, T. M.,
Peterson, K., Pierre, N., Pisan, C., Pollara, V., Raymond, C.,
Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S.,
Seevery, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Subdramanian, A., Talamas, J., Testa, S., Theodore, J., Tirrell, A.,
Travers, M., Triggilo, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B.,
Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zimmer, A. and
Zody, M.

TITLE Direct Submission
 JOURNAL Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 COMMENT On Jan 13, 2000 this sequence version replaced g1:5970502. All repeats were identified using RepeatMasker:
 Smit, A. F. A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
 ----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@
Project Information

Center project name: L3985
Center clone name: 21_D_18

* NOTE: This record contains 55 individual

* sequencing reads that have not been assembled into contigs. Runs of N are used to separate the reads and the order in which they appear is completely arbitrary. Low-pass sequence sampling is useful for identifying clones that may be gene-rich and allows overlap relationships among clones to be deduced. However, it should not be assumed that this clone will be sequenced to completion. In the event that the record is updated, the accession number will be preserved.		
1	869: contig of 869 bp in length	
870	969: gap of 100 bp	
970	1815: contig of 846 bp in length	
1816	1915: gap of 100 bp	
1916	2797: contig of 882 bp in length	
2798	2897: gap of 100 bp	
2898	3768: contig of 871 bp in length	
3769	3868: gap of 100 bp	
3869	4769: contig of 901 bp in length	
4770	4869: gap of 100 bp	
4870	5739: contig of 870 bp in length	
5740	5839: gap of 100 bp	
5840	6703: contig of 864 bp in length	
6704	6803: gap of 100 bp	
6804	7675: contig of 872 bp in length	
7676	7775: gap of 100 bp	
7776	8671: contig of 896 bp in length	
8672	8771: gap of 100 bp	
8772	9639: contig of 868 bp in length	
9640	9739: gap of 100 bp	
9740	10613: contig of 874 bp in length	
10614	10713: gap of 100 bp	
10714	11572: contig of 859 bp in length	
11573	11672: gap of 100 bp	
11673	12528: contig of 856 bp in length	
12529	12628: gap of 100 bp	
12629	13488: contig of 860 bp in length	
13489	13588: gap of 100 bp	
13589	14469: contig of 881 bp in length	
14470	14569: gap of 100 bp	
14570	15452: contig of 883 bp in length	
15453	15552: gap of 100 bp	
15553	16429: contig of 877 bp in length	
16430	16529: gap of 100 bp	
16530	17401: contig of 872 bp in length	
17402	17501: gap of 100 bp	
17502	18387: contig of 886 bp in length	
18388	18487: gap of 100 bp	
18488	19361: contig of 874 bp in length	
19362	19461: gap of 100 bp	
19462	20365: contig of 865 bp in length	
20327	20426: gap of 100 bp	
20427	21291: contig of 865 bp in length	
21292	21391: gap of 100 bp	
21392	22261: contig of 890 bp in length	
22282	22381: gap of 100 bp	
22382	23271: contig of 890 bp in length	
23272	23371: gap of 100 bp	
23372	24249: contig of 878 bp in length	
24250	24349: gap of 100 bp	
24350	25213: contig of 864 bp in length	
25214	25313: gap of 100 bp	
25314	26162: contig of 849 bp in length	
26163	26262: gap of 100 bp	
26263	27122: contig of 860 bp in length	
27123	27222: gap of 100 bp	
27223	28006: contig of 864 bp in length	
28087	28186: gap of 100 bp	
28187	29073: contig of 887 bp in length	
29074	29173: gap of 100 bp	
29174	30055: contig of 882 bp in length	
30056	30155: gap of 100 bp	
30156	31136: contig of 981 bp in length	

[illegible]

	DB	134042	AATTAATAAAAAAAAAAAAAAATATATAAATTAATAATAAAAAAAAAATGAATTAAATAAA	154101
Oy	241	taaaaaaaattccgagggggaataaataatgaaaaaaatattcaagttactgtatc	300	
Db	154102	TAAAAATTTATATATATATATATATATATATATATATATATATATATGCAATTTATTCACA	154161	
Oy	301	ttaagtcttcacgtcttctcatgct	327	
Db	154162	TTTTAGTGTGGTGTTTTCCTTTATTT	154188	
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	DEFINITION	Plasmodium falciparum chromosome 12 clone 3D7, ** SEQUENCING IN PROGRESS **; 4 unordered pieces.		
	ACCESSION	AC005506		
	VERSION	AC005506.8	GI:9797717	
	KEYWORDS	HTG: HTGS_PHASE1.		
	SOURCE	Malaria parasite P. falciparum.		
	ORGANISM	Plasmodium falciparum		
	REFERENCE	Eukaryote; Alveolata; Apicomplexa; Haemosporida; Plasmodium.		
	AUTHORS	1 (bases 1 to 205429)		
		Hyman,R.W., Fung,E.L., Qin,F., Rowley,D., Mao,J., Tamaki,T.,		
		Kundi,O.B., Conway,A.B. and Davis,R.W.		
	TITLE	Plasmodium falciparum 3D7 chromosome 12		
	JOURNAL	Unpublished		
	REFERENCE	2 (bases 1 to 205429)		
	AUTHORS	Hyman,R.W., Qin,F., Fung,E.L., Conway,A.B. and Davis,R.W.		
	JOURNAL	Direct Submission		
		Submitted (21-AUG-1998) Stanford DNA Sequencing and Technology Center, Stanford University, 855 California Avenue, Palo Alto, CA 94304, USA		
	COMMENT	On Aug 12, 2000 this sequence version replaced gi:8810449. * NOTE: This is a 'working draft' sequence. It currently * consists of 4 contigs. The true order of the pieces * is not known and their order in this sequence record is * arbitrary. Gaps between the contigs are represented as * runs of N, but the exact sizes of the gaps are unknown. * This record will be updated with the finished sequence * as soon as it is available and the accession number will * be preserved.		
		* 1 132269: contig of 132269 bp in length		
		* 132270 132469: gap of unknown length		
		* 132470 192227: contig of 59758 bp in length		
		* 192228 192427: gap of unknown length		
		* 192428 203864: contig of 11437 bp in length		
		* 203865 204064: gap of unknown length		
		* 204065 205429: contig of 1365 bp in length.		
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		/chromosome="12"		
		/clone="PFYAC357"		
		/clone="3D7"		
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	Best Local Similarity	52.3%; Pred. No. 0.49;		
	Matches 171, Conservative	0; Mismatches 156; Indels 0; Gaps		
Oy	1	attgggatattctaataatttagcacgaagaatglttaaatgaaataagaataaaaa	60	
Db	133530	ATTTAGAGGATTTTTATTGAGCCCGTCGCCGTAGGCAAAAAAAAAAAATATAT	133589	
Oy	61	gatatattaattatagtcgaaaattcataattatagtagtaagtagttaataataa	120	
Db	133590	AATTAATAATAATAATAAAAAAAAAAAATTTTATTATATAAATAATTAATAAAAAA	133649	

Oy	121	aaaggttcctcgggggacacatttctgtttcaaaaaagaaatacataaattgcat	180
Db	133650	AAAAAAAAATTATTAATTAAATTAAATTAATAAAAAAAAAAAAAAAAAAAATATATATAATA	133709
Oy	181	aaagtcgaataataattatcttattttaaatttgtaaaaaattgataataattg	240
Db	133710	AATTAATATATAAAAAAAAAAATATATATAATAATAATAATAATAAATGAATAATAATA	133769
Oy	241	taaaaaaaattcagggggaataataatgaaaaaaataattcagaagttactgtaatt	300
Db	133770	TAAAAAT	133829
Oy	301	ttatgtttcatgcttcttactgtt	327
Db	133830	TTTATGTGTGTGTGTGTTCCTTTATAT	133856
RESULT	10		
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DEFINITION	AF300334	3392 bp	DNA
ACCESSION	AF300334		INV
VERSION	AF300334		12-NOV-2000
KEYWORDS	AF300334.1	GI:11139097	
SOURCE			
ORGANISM			
REFERENCE			
AUTHORS			
TITLE			
JOURNAL			
REFERENCE			
AUTHORS			
TITLE			
JOURNAL			
FEATURES			
source			
gene			
promoter			
mRNA			
CDS			
BASE COUNT			
ORIGIN			
Query Match			
Best Local Similarity			
Matches			
Oy	11	tcttaaatcttgacacagagaagtgtttaatgaataaagataaagaataaagatatataa	70
Db	1875	TTTTGATATTACTTTGATATATTTTTTTTAATTAATTAATAAAAAAAAAAAAAAAAAACATA	1816

QY	71	ttatctagctgaacatttataatctatgaatgaatcatgattctataataaagaagtcttct	130
Db	1815	ATATTTTTGGAAAAAATAAATTTATTATAATGGAATTAATATAATTAATAATCA	1756
QY	131	cgggggacacttttctgttttaaaagggaataataaataatctagaataaagctga	190
Db	1755	ATTGATGCTGCTGCTGAATGAATAATATGATTTTATTATTAATGATTCATTATTAT	1696
QY	191	aataatcttttcttataatcttataatcttgaataatctgataatctgaatctga	250
Db	1695	TATATATTTATTTATTTTAAAAATTTTCAAAATTT-TTTTAAATAAATATAAATAA	1637
QY	251	ttcaggggggaataataatgaataaataatcttcaaaagttctacgttaattctgt	310
Db	1636	TTTACTGTTTATCTATATTAATTAATTAATGTTTATATTAATTTTATTTTATTT	1577
QY	311	catgttcttatttgtt	327
Db	1576	TTTTTTTTTTTATTTTTT	1560
RESULT	11		
AC079314/c			
LOCUS			
DEFINITION	AC079314 242513 bp DNA HTG 09-AUG-2001		
SEQUENCE	28 unordered pieces.		
ACCESSION	AC079314		
VERSION	AC079314.25 GI:15042762		
KEYWORDS	HTG; HTGS-PHASE1; HTGS-DRAFT.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo. 1 (bases 1 to 242513)		
AUTHORS	Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C., Alsbrooks,S.L., Amaraltinge,H.C., Are,J.R., Banks,T., Barbara,J., Benton,J., Blamege,K., Blankenburg,K., Bonnin,D., Bouch,J., Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Bunay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,L., Chowdry,I., Christopoulos,C., Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C., Day-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K., Harris,C., Harris,K., Hart,M., Haylak,P., Hawes,A., Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C., Hollins,B., Homs,F., Howard,S., Huber,J., Hulys,S., Hume,J., Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudan,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J., Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L., Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W., Louisgied,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E., Massey,E., Kawhney,E., Mcleod,M.P., Meador,M., Mel,G., Metzker,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newson,N., Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nomenkwo,S., Oguh,M., Okunoye,G., Orangunye,N., Oviedo,R., Pace,A., Payton,B., Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L., Quiles,M., Ren,Y., Rives,M., Rojs,A., Rojibokan,I., Rote,M., Ruiz,S., Saevry,G., Scherer,S., Scott,G., Shen,H., Shooshlati,N., Slusson,I., Sodergren,E., Sonalke,T., Sparks,A., Stanley,H., Stone,H., Sultan,A., Svatek,A., Taber,P., Tamerlisa,A., Tamerlisa,K., Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S., Usmani,K., Vasquez,L., Vera,Y., Villalob,D., Vinson,R., Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S., Williams,G., Williamson,A., Wlezyk,R., Wooden,S., Wortley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorilla,S., Nelson,D.,		

REFERENCE Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.

1 (bases 1 to 318221)

AUTHORS Bowman, S., Churcher, C., Harris, B., Harris, D., Lawson, D., Quail, M.

TITLE Direct Submission

JOURNAL Submitted (15-MAR-1999) P.falciparum Genome Sequencing Consortium, The Sanger Centre, Wellcome Trust Genome Camps, Hinxton, Cambridge CB10 1SA, UK

COMMENT On Aug 24, 1999 this sequence version replaced g1:573182.

For more information about this sequence or the Malaria Project, see <http://www.sanger.ac.uk/Projects/P.falciparum>. IMPORTANT: This sequence is unfinished and does not necessarily represent the correct sequence. Work on the sequence is in progress and the release of this data is based on the understanding that the sequence may change as work continues. The sequence may be contaminated with foreign sequence from E.coli, yeast, vector, phage etc.

* NOTE: This is a 'working draft' sequence.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

FEATURES Location/Qualifiers

1..318221

/organism="Plasmodium falciparum"

/strain="3D7"

/db_xref="taxon:5833"

/chromosome="13"

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ORIGIN

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Best Local Similarity 52.08; Pred. No. 0.69; Mismatches 157; Indels 0; Gaps 0;

Matches 170; Conservative 0; Mismatches 157; Indels 0; Gaps 0;

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Db 141159 ATTGTTTCTTTTAAATATATACATAGAAATTTTTCATTAATAATATACATTA 141100

QY 61 gataatataatagctgaaattatataatgataagataagtaataataa 120

Db 141099 AATTAATATATAGATATATATATATATATATATATATATATATATAT 141040

QY 121 aaagtgctcggggggaacttttctttaaaggaaataataataaattgat 180

Db 141039 AATGCTGCAATATATATATATATATATATATATATATATATATATAT 140980

QY 181 aaagtgtaaaataatttttactttaaattgtaaaattgataatgaattg 240

Db 140979 AAT 140920

QY 241 taaataaaattcaggggggaataaaatgaataaattatcaagttactgtaatt 300

Db 140919 ATATCTAT 140860

QY 301 ttatgtttcaatgctttcttctatgctt 327

Db 140859 ATATTTTGAATTTCTTTTGTTTT 140833

RESULT 13

AF377947/c

LOCUS

DEFINITION

PROGRESS ***

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

AF377947 140414 bp DNA HTG 15-MAY-2001

Oryza sativa chromosome 3 clone OSUNBA003ZE21, *** SEQUENCING IN

PROGRESS ***

AF377947

HTG: HTGS PHASE1.

Oryza sativa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

REFERENCE Eukaryota; Oryzaeae; Oryza.

1 (bases 1 to 140414)

AUTHORS Eastman, A.P., Smith, S.C., Gingle, A., Pratt, L.H. and

TITLE Direct Submission

JOURNAL Submitted (01-MAY-2001) Botany, University of Georgia, Miller Plant Sciences, Athens, Georgia 30602, USA

COMMENT * NOTE: This is a 'working draft' sequence. It currently

* consists of 5 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

1..140414

/organism="Oryza sativa"

/db_xref="taxon:4530"

/chromosome="3"

/clone="OSUNBA003ZE21"

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ORIGIN

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Best Local Similarity 55.88; Pred. No. 0.83; Mismatches 163; Conservative 0; Mismatches 126; Indels 3; Gaps 1;

Matches 163; Conservative 0; Mismatches 126; Indels 3; Gaps 1;

QY 35 tttaaatgaataaagataaagaatataatataatagctgaataattat 94

Db 21415 TTAAATTAATTTTAAATATATATATATATATATATATATATATATATAT 21356

QY 95 atatgaatagatagtttaataaataaagctgctcggggggaact---tttggttt 151

Db 21355 ATTAATAAATAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 21296

QY 152 aaaaaggaataataaataaattagataaagtgtaataataatttttacttaaa 211

Db 21295 TATTAATAAATAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 21236

QY 212 ttgttaaaatttgatataattgtaataaataaattcaggggggaataata 271

Db 21235 TAAATTTTAAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 21176

QY 272 aaaaattatcagaagttactgtaattttatgcttcaatgcttctat 323

Db 21175 ATATATATATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 21124

RESULT 14

AC068139/c

LOCUS

DEFINITION

PROGRESS ***

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

human.

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 178783)

Smit, D.R.

Genome Therapeutics Corporation Sequencing Center; Human Genome

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:10:08 ; Search time 2272.52 Seconds
(without alignments)
1546.244 Million cell updates/sec

Title: US-09-531-438-3

Sequence: 1 atttggatcttcttaattt.....tttcattgttcttattgtt 327

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapept 1.0

Searched: 11351937 seqs, 5372889281 residues

Total number of hits satisfying chosen parameters: 22703874

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST: *
1: em_estfun: *
2: em_esthum: *
3: em_estin: *
4: em_estom: *
5: em_estpl: *
6: em_estba: *
7: em_estro: *
8: em_estov: *
9: em_hic: *
10: gb_estl: *
11: gb_est2: *
12: gb_hic: *
13: gb_gss: *
14: em_gss_fun: *
15: em_gss_hum: *
16: em_gss_inv: *
17: em_gss_pln: *
18: em_gss_pro: *
19: em_gss_rtd: *
20: em_gss_vrt: *
21: em_gss_other: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	91	27.8	1101	13	AL069706 Drosophila
2	89.2	27.3	1225	13	AL106171 Drosophila
3	85.4	26.1	1101	13	AL064091 Drosophila
4	84.6	25.9	1101	13	AL063921 Drosophila
5	84.4	25.8	905	13	AL077798 Drosophila
6	83.2	25.4	524	13	AL07541 Tetradon
7	83	25.4	1101	13	AL052544 Drosophila
8	83	25.4	1101	13	AL069440 Drosophila
9	82.8	25.3	987	13	AL104456 Drosophila
10	82.2	25.1	1101	13	AL060732 Drosophila
11	81.2	24.8	734	13	AL099163 Drosophila
12	81.2	24.8	1200	13	AL106578 Drosophila

C 13	81	24.8	1125	10	AL547503	AL547503 Tetradon
C 14	80.6	24.6	1092	13	CNS020K7	AL175696 Tetradon
C 15	80.4	24.6	973	13	CNS0080F	AL052232 Drosophila
C 16	80.4	24.6	1001	13	CNS0155H	AL105023 Drosophila
C 17	80.4	24.6	1101	13	CNS00FMC	AL070972 Drosophila
C 18	80.2	24.5	836	13	CNS02M02	AL217379 Tetradon
C 19	79.6	24.3	694	13	AO853360	AO853360 nbxb0003G
C 20	79.4	24.3	1101	13	CNS00EVL	AL069706 Drosophila
C 21	79	24.2	966	13	CNS0052C	AL061991 Drosophila
C 22	79	24.2	1101	13	CNS0021J	AL061936 Drosophila
C 23	78.6	24.0	581	13	CNS034DK	AL227297 Tetradon
C 24	78.6	24.0	928	13	CNS00DXY	AL071865 Drosophila
C 25	78.6	24.0	996	13	CNS00FJH	AL071063 Drosophila
C 26	78.4	24.0	1069	13	CNS0107G	AL098614 Drosophila
C 27	78.4	24.0	1101	13	CNS00E07	AL069440 Drosophila
C 28	78.4	24.0	1101	13	CNS016L1	AL106896 Drosophila
C 29	78	23.9	876	13	CNS0096L	AL053529 Drosophila
C 30	77.8	23.8	987	13	CNS014P0	AL104456 Drosophila
C 31	77.6	23.7	1101	13	CNS00EYX	AL071206 Drosophila
C 32	77.6	23.7	1101	13	CNS00K28	AL097166 Drosophila
C 33	77.4	23.7	770	13	AO740708	AO740708 HS_5307_A
C 34	77.4	23.7	1101	13	CNS000B8	AL063632 Drosophila
C 35	77.4	23.7	1101	13	CNS00EPO	AL069493 Drosophila
C 36	77.4	23.7	1225	13	CNS0161D	AL106171 Drosophila
C 37	77.2	23.6	1101	13	CNS00KAE	AL077628 Drosophila
C 38	77.2	23.6	1200	13	CNS016C0	AL106578 Drosophila
C 39	76.8	23.5	614	13	CNS0152H	AL104915 Drosophila
C 40	76.8	23.5	1099	10	AL536986	AL536986 AL536986
C 41	76.6	23.4	1101	13	CNS0039Q	AL063931 Drosophila
C 42	76.4	23.4	1092	13	CNS020K7	AL175696 Tetradon
C 43	76.2	23.3	928	13	CNS00DKY	AL071865 Drosophila
C 44	76.2	23.3	1203	13	CNS015MU	AL106008 Drosophila
C 45	76	23.2	1124	13	CNS073BM	AL427304 clone BAO

ALIGNMENTS

RESULT 1
CNS00EVL
LOCUS
DEFINITION
BACR29B23 of Rpci-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Drosophila melanogaster
fruit fly.
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osoegawa and Aaron Mammoler in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named Rpci-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's pl and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

```

FEATURES
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    /db_xref="taxon:7227"
    /clone_id="RPCI-98"
    /clone="BACR29B23"
    /note="end : 17"

BASE COUNT
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ORIGIN

Query Match
Best Local Similarity 27.8%; Score 91; DB 13; Length 1101;
Matches 127; Conservative 68; Mismatches 108; Indels 2; Gaps 1

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Db 700 AAAAAAAMATWAAATWATWATWATWATWATWATWATWATWATWATWATWATWATWATW 759

OY 75 atagctgaagaattcaatcatatgataagtaagtaagtaataaagaattcttcg9g 134
    || : : || : || : || : || : || : || : || : || : || : || : || :
Db 760 ATATATATWTTWATWATWATWATWATWATWATWATWATWATWATWATWATWATWATW 819

OY 135 ggaacattttgtc--ttaaagggaataataaataalttgataaagctgtaaaa 192
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OY 193 taattatttttaatttaattgtttaaagaatttgatcatatgaatttgtaaaaaaatt 252
    || : : || : || : || : || : || : || : || : || : || : || : || :
Db 880 TAAATTTTWTWTTTWTWATWATWATWATWATWATWATWATWATWATWATWATWATW 939

OY 253 caggggggaataataaataaataattcaaaagttactcgttaattttatgtttca 312
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Db 940 WATATTWTTTATTAAMWTATWATWATWATWATWATWATWATWATWATWATWATWATW 999

OY 313 tgttt 317
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Db 1000 TAAAT 1004

RESULT 2
LOCUS CNS0161D/c
DEFINITION Drosophila melanogaster genome survey sequence Sp6 end of BAC
          BACN1518 of DrosBAC library from Drosophila melanogaster (fruit
          fly), genomic survey sequence.
ACCESSION AL106171
VERSION AL106171.1 GI:5620504
KEYWORDS GSS.
SOURCE fruit fly
          Plasmid Drosophila melanogaster
          Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
          Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
          Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
          1 (bases 1 to 1225)
          Genoscope.
          Direct Submission
          Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage
          BP 191 91006 EVRY cedex - FRANCE (E-mail : segre@genoscope.cns.fr
          - Web : www.genoscope.cns.fr)
          Determination of this BAC-end sequence was carried out as part of a
          collaboration with the European Drosophila Genome Project (EDGP) -
          http://www.edgp.ebi.ac.uk/. This Drosophila melanogaster BAC
          library (Dros BAC) was made by Alain Billard at CEPH (Centre
          d'Etude du Polymorphisme Humain) with funding provided by a MRC
          project grant. The DNA was prepared from embryos by Alain Bucheton
          and Genevieve Payan. It has been constructed in the vector
          pbeloBAC11.

FEATURES
source
    Location/Qualifiers
    1. 1225
    /organism="Drosophila melanogaster"
    /plasmid="pbeloBAC11"
    /db_xref="taxon:7227"

```

			/clone.lib="DrosBAC"	
			/clone="BACN15C18"	
			/note="end : SP6"	
BASE COUNT	266 a	128 c	38 g	368 t 425 others
ORIGIN				
Query Match	27.3%	Score 89.2:	DB 13;	Length 1225;
Best Local Similarity	40.8%;	Pred. No. 0.014;		
Matches 125;	Conservative 61;	Mismatches 120;	Indels 0;	Gaps 0;
Oy	8	aatacttaattgcacagaagatgctttaactgaataagaataaaagatatat 67		
Db	1164	ATATAATWTAATTAAWWMAAAMWTTTTATATATATWTTATATATAAAAATAATATMM 1105		
Oy	68	taataataatgcgtcaaatcataatcatatgatagaagtatcgtaaataaaaagtcg 127		
Db	1104	WAWWWAAAAAAMWAAMAAWMTWTTWRRAATATAAAATATATATATATAAATAWAMMW 1045		
Oy	128	tctcggggagaccttttcttgctttaaaaggaaataataaattagataaaagtg 187		
Db	1044	TWMAAAATATATTTTTTTTTTTTWTATNAAAAAAAAAAAAAATATATATWATRAAAMW 985		
Oy	188	taaaataatcttatctttaaattgltaaaaaattgataaattgaaattgaaanaa 247		
Db	984	TAAAAAAAAATAATATWATWMAWTTTTTTTRAAAMWTTTTTTTTTTWTTAAAMWT 925		
Oy	248	aattccaggggggaataaaatgaaaataattattccaagttcacgttaatttaagt 307		
Db	924	WWTATWTTTWTWTRRRARAATWMTTTTTTTTTTTTTTTTATATTTTTTTTMMHHT 865		
Oy	308	ttcat 313		
Db	864	XYMYT 859		
RESULT 3				
CNS003BP/c				
LOCUS	CNS003BD	1101 bp	DNA	GSS 03-JUN-1999
DEFINITION	Drosophila melanogaster genome survey sequence TET3 end of BAC # BACR08K08 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.			
ACCESSION	AI064091			
VERSION	AI064091.1	GI:4941847		
KEYWORDS	GSS.			
ORGANISM	fruit fly ;			
SOURCE	Drosophila melanogaster			
REFERENCE	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.			
AUTHORS	I (bases 1 to 1101)			
JOURNAL	Genoscope.			
COMMENT	Direct Submission Submitted (02-JUN-1999) Genoscope - Centre National de Sequenage : BP 191 91006 EVRY cedex - FRANCE (E-mail : segr@genoscope.cns.fr) Web : www.genoscope.cns.fr) Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see http://www.fruitfly.org The BDGP Drosophila melanogaster BAC library was prepared by Kazuroyo Osoegawa and Aaron Mamoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's pl and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm. Location/Qualifiers 1..1101			
FEATURES				
SOURCE				

Genoscope. Direct Submission
Submitted (02-JUN-1999) Genoscope - Centre National de Séquençage :
EP 131 91006 EVRY cedex - FRANCE (E-mail : seqre@genoscope.cns.fr
- web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see <http://www.fruitfly.org/TheBDGP/Drosophila>
melanogaster BAC library was prepared by Kazuhiro Oseogawa and
Aron Memmoser in Pieret de Jong's laboratory in the Department of
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
NY. The library is named RPc1-98 and was constructed by partial
EcoRI digestion of Drosophila DNA provided by the BDGP from the
isogenic strain y2; cn bw sp, the same strain used for the BDGP's
c1 and EST libraries. A more detailed description of the library
and how to order individual BAC clones, the entire library, or

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:50:52 ; Search time 216.42 Seconds
(Without alignments)
1295.376 Million cell updates/sec

Title: US-09-531-438-3
Perfect score: 327
Sequence: 1 atttgagatcttaattt.....tttcattgtttctattgt 327

Scoring table: IDENTITY_NIC
Gapop 10.0 , Gapept 1.0

Searched: 930621 seqs, 428662619 residues
Total number of hits satisfying chosen parameters: 1861242

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: /SID52/gcgdata/geneseq/NA1980.DAT:*
2: /SID52/gcgdata/geneseq/NA1981.DAT:*
3: /SID52/gcgdata/geneseq/NA1982.DAT:*
4: /SID52/gcgdata/geneseq/NA1983.DAT:*
5: /SID52/gcgdata/geneseq/NA1984.DAT:*
6: /SID52/gcgdata/geneseq/NA1985.DAT:*
7: /SID52/gcgdata/geneseq/NA1986.DAT:*
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22: /SID52/gcgdata/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	327	100.0	327	20	AAK60300
2	327	100.0	1392	20	AAK60299
3	130.2	39.8	22	AAK60299	Promoter of the b
4	130.2	39.8	22	AAK60299	DNA encoding the b
5	130.2	39.8	22	AAK60299	Oligonucleotide D1
6	130.2	39.8	22	AAK60299	Oligonucleotide D1
7	130.2	39.8	22	AAK60299	Oligonucleotide D1
8	130.2	39.8	22	AAK60299	Oligonucleotide D2
9	129.8	39.7	22	AAK60299	Oligonucleotide D1
10	129.8	39.7	22	AAK60299	Oligonucleotide D1
11	129.8	39.7	22	AAK60299	Oligonucleotide D1

C 12	129.8	39.7	936	22	AAK60299
C 13	129.8	39.7	936	22	AAK60299
C 14	129.8	39.7	936	22	AAK60299
C 15	91.6	28.0	244	22	AAK60299
C 16	90.8	27.8	244	22	AAK60299
C 17	66.8	20.4	4590	22	AAK60299
C 18	66.4	20.3	2486	21	AAK60299
C 19	64.8	19.5	20674	21	AAK60299
C 20	63.8	19.3	2435	21	AAK60299
C 21	63.2	19.0	9789	17	AAK60299
C 22	62.2	18.8	6243	20	AAK60299
C 23	61.6	18.8	19124	18	AAK60299
C 24	61.6	18.8	19124	21	AAK60299
C 25	61	18.7	700	22	AAK60299
C 26	61	18.7	5852	12	AAK60299
C 27	60.4	18.5	700	22	AAK60299
C 28	60.2	18.4	1341	20	AAK60299
C 29	60.2	18.4	1410	20	AAK60299
C 30	59.6	18.2	513445	22	AAK60299
C 31	59.4	18.2	6265	20	AAK60299
C 32	59	18.0	381	22	AAK60299
C 33	58.8	18.0	53585	20	AAK60299
C 34	58.4	17.9	3975	9	AAK60299
C 35	58.4	17.9	3975	13	AAK60299
C 36	58.4	17.9	163319	21	AAK60299
C 37	58.2	17.8	1132	21	AAK60299
C 38	58.2	17.8	2503	15	AAK60299
C 39	58.2	17.8	9048	18	AAK60299
C 40	57.8	17.7	8310	20	AAK60299
C 41	57.6	17.6	1907	20	AAK60299
C 42	57.4	17.6	366	22	AAK60299
C 43	57	17.4	20674	21	AAK60299
C 44	56.8	17.4	605	17	AAK60299
C 45	56.8	17.4	665	21	AAK60299

ALIGNMENTS

RESULT 1	AAK60300 standard; DNA; 327 BP.
ID	AAK60300
AC	AAK60300;
XX	12-AUG-1999 (first entry)
XX	Promoter of the beta-2 toxin gene of Clostridium perfringens type C.
XX	Beta-2 toxin; Clostridium perfringens type C; gene promoter;
XX	vaccine; Clostridium tetani; ss.
XX	Clostridium perfringens.
XX	FR2768747-A1.
XX	26-MAR-1999.
XX	19-SEP-1997; 97FR-0011710.
XX	19-SEP-1997; 97FR-0011710.
XX	(INSP) INST PASTEUR.
XX	Gilbert M, Popoff MR;
XX	WPI; 1999-217498/19.
XX	Clostridium beta2 toxin gene promoter and signal sequence - useful
XX	against toxins from Clostridium perfringens
XX	Claim 1; Page 32; 46pp; French.

CC The present sequence represents the promoter of the beta-2 toxin
CC gene of Clostridium perfringens type C. The beta2-toxin promoter
CC and gene sequences can be used to produce vaccines against Clostridium,
CC and especially Clostridium perfringens, or Clostridium
CC tetani.

CC Sequence 327 BP; 141 A; 13 C; 44 G; 129 T; 0 other;

Query Match 100.0%; Score 327; DB 20; Length 327;

Best Local Similarity 100.0%; Pred. No. 2.2e-35;

Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atttggatattttaaatttagcacagaagaatgtttaaatgaataaataa 60
DB 1 atttggatattttaaatttagcacagaagaatgtttaaatgaataaataa 60
QY 61 gatataattatataagctggaattataattatataagtagtaataataa 120
DB 61 gatataattatataagctggaattataattatataagtagtaataataa 120
QY 121 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 180
DB 121 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 180
QY 181 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 240
DB 181 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 240
QY 241 taaaaaaatttcagggggaataataatgaataaataatttcaagttactgtaatt 300
DB 241 taaaaaaatttcagggggaataataatgaataaataatttcaagttactgtaatt 300
QY 301 ttatgttttcattgttttttcttattgtt 327
DB 301 ttatgttttcattgttttttcttattgtt 327

RESULT 2

AAK60299 standard; DNA; 1392 BP.

AAK60299;

12-AUG-1999 (first entry)

DNA encoding the beta-2 toxin of Clostridium perfringens type C.

Beta-2 toxin; Clostridium perfringens type C; gene promoter;
KW vaccine; Clostridium tetani; ss.

Clostridium perfringens.

FR2768747-A1.

26-MAR-1999.

19-SEP-1997; 97FR-0011710.

19-SEP-1997; 97FR-0011710.

(INSP) INST PASTEUR.

Gibert M, Popoff MR.

WPI, 1999-217498/19.

P-PSDB; AAY16591.

Clostridium beta2 toxin gene promoter and signal sequence - useful
PT against toxins from Clostridium perfringens

Example A; Page 31; 46pp; French.

CC The present sequence encodes the beta-2 toxin of Clostridium
CC perfringens type C. The specification describes the Clostridium
CC perfringens beta 2 toxin gene promoter (see AAK60300). The
CC sequences can be used to produce vaccines against Clostridium,
CC and especially Clostridium perfringens, or Clostridium
CC tetani.

CC Sequence 1392 BP; 606 A; 115 C; 209 G; 462 T; 0 other;

Query Match 100.0%; Score 327; DB 20; Length 1392;

Best Local Similarity 100.0%; Pred. No. 1.7e-35;

Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atttggatattttaaatttagcacagaagaatgtttaaatgaataaataa 60
DB 1 atttggatattttaaatttagcacagaagaatgtttaaatgaataaataa 60
QY 61 gatataattatataagctggaattataattatataagtagtaataataa 120
DB 61 gatataattatataagctggaattataattatataagtagtaataataa 120
QY 121 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 180
DB 121 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 180
QY 181 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 240
DB 181 aaagtgttcgaggacacattttgttttaaaaggaataataaataattagat 240
QY 241 taaaaaaatttcagggggaataataatgaataaataatttcaagttactgtaatt 300
DB 241 taaaaaaatttcagggggaataataatgaataaataatttcaagttactgtaatt 300
QY 301 ttatgttttcattgttttttcttattgtt 327
DB 301 ttatgttttcattgttttttcttattgtt 327

RESULT 3

AAF58252 standard; DNA; 936 BP.

AAF58252;

24-APR-2001 (first entry)

Oligonucleotide D1835.

Electron-transfer group; ETM; mismatch; genotyping;
KW gene expression; ss.

Synthetic.

WO200107665-A2.

01-FEB-2001.

26-JUL-2000; 2000WO-US20476.

26-JUL-1999; 99US-0145695.

17-MAR-2000; 2000US-0190259.

(CLIN-) CLINICAL MICRO SENSORS INC.

umek RM;

WPI, 2001-159728/16.

Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface

PI Umek RM;
 XX WPI: 2001-159728/16.
 XX Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface -
 XX
 PS Example 6; Page 128; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 CC
 XX Sequence 936 BP; 5 A; 139 C; 10 G; 6 T; 776 other;

Query Match 39.8%; Score 130.2; DB 22; Length 936;
 Best Local Similarity 0.9%; Pred. No. 9.2e-10;
 Matches 3; Conservative 268; Mismatches 56; Indels 0; Gaps 0;
 QY 1 attgggatacctaatttagcacagaagaatgttaaatgaataaagataaataa 60
 DB 136 www..... 195
 QY 61 gatataataatataagctgaataattataatataatgaataagataatataa 120
 DB 196 gwww..... 255
 QY 121 aaaggtctcgggggacacttttgtttaaaaggaataataaattagat 180
 DB 256 www..... 315
 QY 181 aaagtgtaataataatttttaatttaattgttaaaattgataataatgattg 240
 DB 316 www..... 375
 QY 241 taaaaaaatttcagggggaataataatgaataaattatccaagttcactgaatt 300
 DB 376 www..... 435
 QY 301 ttatgtttcatgttttctatgtt 327
 DB 436 www..... 462

RESULT 8
 AAF58255
 ID AAF58255 standard; DNA; 938 BP.
 XX
 AC AAF58255;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D1876.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 XX
 OS Synthetic.
 XX
 PN WO200107665-A2.
 XX
 PD 01-FEB-2001.
 XX
 PF 26-JUL-2000; 2000WO-US20476.
 XX
 PR 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX

PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 PI Umek RM;
 XX WPI: 2001-159728/16.
 XX Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface -
 XX
 PS Example 6; Page 127; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 CC
 XX Sequence 938 BP; 4 A; 144 C; 9 G; 5 T; 776 other;

Query Match 39.8%; Score 130.2; DB 22; Length 938;
 Best Local Similarity 0.9%; Pred. No. 9.2e-10;
 Matches 3; Conservative 268; Mismatches 56; Indels 0; Gaps 0;
 QY 1 attgggatacctaatttagcacagaagaatgttaaatgaataaagataaataa 60
 DB 136 www..... 195
 QY 61 gatataataatataagctgaataattataatataatgaataagataatataa 120
 DB 196 gwww..... 255
 QY 121 aaaggtctcgggggacacttttgtttaaaaggaataataaattagat 180
 DB 256 www..... 315
 QY 181 aaagtgtaataataatttttaatttaattgttaaaattgataataatgattg 240
 DB 316 www..... 375
 QY 241 taaaaaaatttcagggggaataataatgaataaattatccaagttcactgaatt 300
 DB 376 www..... 435
 QY 301 ttatgtttcatgttttctatgtt 327
 DB 436 www..... 462

RESULT 9
 AAF58252/c
 ID AAF58252 standard; DNA; 936 BP.
 XX
 AC AAF58252;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D1835.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 XX
 OS Synthetic.
 XX
 PN WO200107665-A2.
 XX
 PD 01-FEB-2001.
 XX
 PF 26-JUL-2000; 2000WO-US20476.
 XX
 PR 26-JUL-1999; 99US-0145695.
 XX

PR 17-MAR-2000; 2000US-0190259.
XX
PA (CLIN-) CLINICAL MICRO SENSORS INC.
XX
PI
PI Umek RM;
XX
DR WPI; 2001-159728/16.
XX
PT Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface -
XX
XX
PS Example 6; Page 127; 159pp; English.
XX
XX The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
CC monitoring gene expression.
XX
XX Sequence 936 BP; 4 A; 139 C; 10 G; 7 T; 776 other;

[illegible]

RESULT	10
AAF58254/c	
ID	AAF58254 standard; DNA; 936 BP.
XX	
AC	AAF58254;
XX	
DT	24-APR-2001 (first entry)
XX	
DE	Oligonucleotide D1875.
XX	
KW	Electron-transfer group; EFM; mismatch; genotyping
KW	gene expression; ss.
XX	
OS	Synthetic.
XX	
PN	W0200107665-A2.
XX	
PD	01-FEB-2001.
XX	
26-JUL-2000;	2000WO-US20476.

XX 26-JUL-1999: 99US-0145695.
PR 17-MAR-2000; 2000US-0190259.
XX
XX (CLIN-) CLINICAL MICRO SENSORS INC.
PA
XX
XX Umek RM;
PI
XX
XX WPI; 2001-159728/16.
DR
XX
XX Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface
XX
XX
XX
XX Example 6; Page 127; 159pp; English.
PS
XX
XX The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
CC monitoring gene expression.

[illegible]

RESULT	11
AAFs8257/c	
ID	AAFs8257 standard, DNA, 936 BP.
XX	
AC	AAFs8257:
XX	
DT	24-Apr-2001 (first entry)
XX	
DE	Oligonucleotide D1954.
XX	
KW	Electron-transfer group; FTM; mismatch; genotyping
KW	gene expression; ss.
XX	
OS	Synthetic.
XX	
PN	W0200107665-A2.
XX	
PD	01-FEB-2001.

XX 26-JUL-2000; 2000MO-US20476.
 PF 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 PI Umek RM;
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface
 PS Example 6; Page 127; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 XX
 SQ Sequence 936 BP; 5 A; 142 C; 7 G; 6 T; 776 other;

Query Match 39.7%; Score 129.8; DB 22; Length 936;
 Best Local Similarity 0.6%; Pred. No. 1e-09;
 Matches 2; Conservative 269; Mismatches 56; Indels 0; Gaps 0;
 Oy 1 attgggatacttaattagcacagaagaatgtttaagaataagaataaaaaa 60
 Db 750 ww 691
 Oy 61 gatacataataatagctgaaattataataatagataagtaataataaa 120
 Db 690 Gww 631
 Oy 121 aaagtctctcggggacacttttctttaaaaagaataataaaattagat 180
 Db 630 Www 571
 Oy 181 aaagtgtcaaaataatttttatttaattgtaaaaaattgataataattg 240
 Db 570 Www 511
 Oy 241 taaaaaaattcagggggaataataatgaaaaaattattcaagtttactg 300
 Db 510 Www 451
 Oy 301 ttatgtttcattgtttcttattgtt 327
 Db 450 Wwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwww 424

RESULT 12
 AAF58259/c
 ID AAF58259 standard; DNA; 936 BP.
 XX
 AC AAF58259;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D2004.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 OS Synthetic.
 XX
 PN WO200107665-A2.

XX 01-FEB-2001.
 PD
 XX 26-JUL-2000; 2000MO-US20476.
 PF 26-JUL-1999; 99US-0145695.
 PR 17-MAR-2000; 2000US-0190259.
 XX
 PA (CLIN-) CLINICAL MICRO SENSORS INC.
 XX
 PI Umek RM;
 DR WPI; 2001-159728/16.
 XX
 PT Nucleic acids containing electron-transfer group, useful as labels in
 PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
 PT a single surface
 PS Example 6; Page 128; 159pp; English.
 XX
 CC The present invention relates to a composition comprising two nucleic
 CC acids each containing an electron-transfer group (ETM) having
 CC different redox potentials. The invention is used for electronic
 CC detection of nucleic acids, especially of substitutions (mismatches)
 CC and single-nucleotide polymorphisms, e.g. for genotyping,
 CC monitoring gene expression.
 XX
 SQ Sequence 936 BP; 6 A; 138 C; 8 G; 8 T; 776 other;

Query Match 39.7%; Score 129.8; DB 22; Length 936;
 Best Local Similarity 0.6%; Pred. No. 1e-09;
 Matches 2; Conservative 269; Mismatches 56; Indels 0; Gaps 0;
 Oy 1 attgggatacttaattagcacagaagaatgtttaagaataagaataaaaa 60
 Db 750 ww 691
 Oy 61 gatacataataatagctgaaattataataatagataagtaataataaa 120
 Db 690 Gww 631
 Oy 121 aaagtctctcggggacacttttctttaaaaagaataataaaattagat 180
 Db 630 Www 571
 Oy 181 aaagtgtcaaaataatttttatttaattgtaaaaaattgataataattg 240
 Db 570 Www 511
 Oy 241 taaaaaaattcagggggaataataatgaaaaaattattcaagtttactg 300
 Db 510 Www 451
 Oy 301 ttatgtttcattgtttcttattgtt 327
 Db 450 Wwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwww 424

RESULT 13
 AAF58262/c
 ID AAF58262 standard; DNA; 936 BP.
 XX
 AC AAF58262;
 XX
 DT 24-APR-2001 (first entry)
 XX
 DE Oligonucleotide D2007.
 XX
 KW Electron-transfer group; ETM; mismatch; genotyping;
 KW gene expression; ss.
 OS Synthetic.

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:51:38 ; Search time 2099.46 Seconds
(without alignments)
1673.702 Million cell updates/sec

Title: US-09-531-438-3
Perfect score: 327
Sequence: 1 atttggatattttaaattt.....tttcattttttatttgtt 327

Scoring table: OLIGO-MNC
Gapop 60.0, Gapext 60.0

Searched: 11351937 seqs, 5372889281 residues

Word size: 0

Total number of hits satisfying chosen parameters: 80718

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

EST:*
1: em_estfun:*
2: em_esthm:*
3: em_estin:*
4: em_estom:*
5: em_estpl:*
6: em_estba:*
7: em_estro:*
8: em_estov:*
9: em_hic:*
10: gb_estl:*
11: gb_est2:*
12: gb_hic:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rnd:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	4.6	42	10	AU011968 AU011968
2	15	4.6	42	10	AU011969 AU011969
3	15	4.6	42	10	AU011971 AU011971
4	15	4.6	42	10	AU011973 AU011973
5	15	4.6	49	10	AA922891 AU011973
6	15	4.6	50	11	AA922891 AU011973
7	15	4.3	24	13	AZ781748
8	14	4.3	28	11	CO1204
9	14	4.3	34	10	AU038857
10	14	4.3	42	10	AW333885
11	14	4.3	42	13	AZ634761
12	14	4.3	46	13	AZ459612

C 13	14	4.3	47	13	AZ345468
14	14	4.3	50	10	BE043289
15	14	4.3	50	13	TA154F10Q
C 16	13	4.0	19	13	AZ331628
C 17	13	4.0	25	13	AZ829725
C 18	13	4.0	26	13	AZ309204
C 19	13	4.0	26	13	AZ866662
C 20	13	4.0	26	13	TA123B12Q
C 21	13	4.0	27	13	AZ784620
C 22	13	4.0	28	13	AZ452653
C 23	13	4.0	30	13	AZ623794
C 24	13	4.0	32	11	H40874
C 25	13	4.0	32	13	AZ458690
C 26	13	4.0	34	10	AA906810
C 27	13	4.0	34	13	AZ586746
C 28	13	4.0	34	13	AZ781725
C 29	13	4.0	35	10	AA246486
C 30	13	4.0	35	11	D45807
C 31	13	4.0	36	13	AZ314238
C 32	13	4.0	37	10	AA913140
C 33	13	4.0	37	10	AU009123
C 34	13	4.0	40	10	AA916625
C 35	13	4.0	40	10	AA922076
C 36	13	4.0	41	11	CO0434
C 37	13	4.0	41	13	AZ777050
C 38	13	4.0	43	13	AZ371136
C 39	13	4.0	43	13	AZ464392
C 40	13	4.0	43	13	AZ575514
C 41	13	4.0	45	11	D20668
C 42	13	4.0	46	13	AZ459612
C 43	13	4.0	46	13	AZ834972
C 44	13	4.0	46	13	AZ991460
C 45	13	4.0	49	10	A1630064

ALIGNMENTS

RESULT 1
AU011968
LOCUS AU011968 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011968 Schizosaccharomyces pombe late log phase cDNA
ACCESSION Schizosaccharomyces pombe cDNA clone spc06162, mRNA sequence.
AU011968
VERSION AU011968.1 GI:3356877
KEYWORDS EST.
SOURCE fission yeast.
ORGANISM Schizosaccharomyces pombe
Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;
Schizosaccharomyces.
REFERENCE 1 (bases 1 to 42)
AUTHORS Moriyo,M. and Mita,K.
TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe
JOURNAL Unpublished (1998)
COMMENT Contact: Mitsuaki Moriyo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-Ku, Chiba 263-8555, Japan
Email: moriyo@nirs.go.jp
FEATURES
source location/Qualifiers
1..42
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06162"
/sex="h minus"
/note="Vector: M13mp19. The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of

Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttttggtttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 2

LOCUS AU011969 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011969 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06163, mRNA sequence.

ACCESSION AU011969
VERSION AU011969.1 GI:3356878

KEYWORDS EST
SOURCE fission yeast.
ORGANISM Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;

REFERENCE 1 (bases 1 to 42)
AUTHORS Morimyo,M. and Mita,K.
TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)
COMMENT Contact: Mitsuoki Morimyo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: morimyo@nirs.go.jp.

FEATURES
source Location/Qualifiers

1..42
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06163"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/sex="h minus"
/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)"

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttttggtttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 3

LOCUS AU011971 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011971 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06165, mRNA sequence.

ACCESSION AU011971
VERSION AU011971.1 GI:3356880
KEYWORDS EST.

SOURCE fission yeast.

ORGANISM Schizosaccharomyces pombe
Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;

REFERENCE 1 (bases 1 to 42)
AUTHORS Morimyo,M. and Mita,K.
TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)

COMMENT Contact: Mitsuoki Morimyo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: morimyo@nirs.go.jp.

FEATURES
source Location/Qualifiers

1..42
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06165"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/sex="h minus"
/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)"

Query Match 4.6%; Score 15; DB 10; Length 42;
Best Local Similarity 100.0%; Pred. No. 9.6e+04;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 142 ttttggtttaaaaa 156
|||||
Db 22 TTTTGTGTTTAAAAA 36

RESULT 4

LOCUS AU011973 42 bp mRNA EST 03-AUG-1998
DEFINITION AU011973 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA clone spc06167, mRNA sequence.

ACCESSION AU011973
VERSION AU011973.1 GI:3356882

KEYWORDS EST
SOURCE fission yeast.
ORGANISM Schizosaccharomyces pombe

Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetales; Schizosaccharomycetaceae;
Schizosaccharomyces.
1 (bases 1 to 42)
AUTHORS Morimyo,M. and Mita,K.
TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe

JOURNAL Unpublished (1998)

COMMENT Contact: Mitsuoki Morimyo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: morimyo@nirs.go.jp.

FEATURES
source Location/Qualifiers

1..42
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc06167"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/sex="h minus"

/note="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <http://www.nirs.go.jp>)"

BASE COUNT 19 a 1 c 7 g 15 t

ORIGIN

Query Match 4.6%; Score 15; DB 10; Length 42;

Best Local Similarity 100.0%; Pred. No. 9.6e+04;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 142 ttttggtttaaaa 156

Db 22 tttttgttttaaaa 36

RESULT 5

AA922891

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

Db 31 AATTGTAAAAAAT 45

RESULT 6

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

SOURCE

FEATURES

SOURCE

FEATURES

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SOURCE

CO1094 50 bp mRNA EST 23-JUL-1996

HUMGSD007754 Human adult (K.Okubo) Homo sapiens cDNA, mRNA

sequence.

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

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CO1094 50 bp mRNA EST 23-JUL-1996

CO1094 50 bp mRNA EST 23-JUL-1996

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert Length: 1000 Std Error: 0.00
Plate: 0021 Row: M Column: 18
Seq primer: CGTGTAAACGACGCGCAGT
Class: Plasmid ends
High quality sequence stop: 24.

FEATURES

source

1. .24

Location/Qualifiers

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone_lib="U0022M0021M18"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g147321419b1AF129072.1), a copy-number inducible derivative of plasmid RL. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

9 a 0 c 3 g 12 t

ORIGIN

Query Match
Best Local Similarity 100.0%; Pred. No. 2.9e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211
|||||
Db 7 ATTATTAATTTTAA 20

RESULT 8
C01204/c 28 bp mRNA EST 23-JUL-1996

LOCUS HUMS0007904 Human adult (K.Okubo) Homo sapiens cDNA, mRNA
DEFINITION
sequence.
C01204
VERSION C01204.1 GI:1433434
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS Okubo, K.
TITLE BodyMap: human gene expression database
JOURNAL Unpublished (1995)
COMMENT Contact: Okubo, K.
Institute for Molecular and Cellular Biol
1-3, Yamada-oka, Suita, Osaka Pref. 565, Japan
Tel: 06-877-5111 (ex. 3315)
Email: kousaku@imcb.osaka-u.ac.jp

Human Gene Signature, 3'-directed cDNA sequence. We are not submitting the same cDNA sequence redundantly to DDBJ since 1993. For the abundance information of clones with this sequence in this library and as well as in other 3'-directed libraries, see <http://www.imcb.osaka-u.ac.jp/bodymap/>. The sequences of the clones represented by this GS sequences is also found there.

FEATURES

source

1. .28

Location/Qualifiers

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone_lib="Human adult (K.Okubo)"

/dev_stage="adult"

/dev_stage="adult"

BASE COUNT

11 a 3 c 3 g 10 t 1 others

ORIGIN

Query Match
Best Local Similarity 100.0%; Pred. No. 2.7e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211
|||||
Db 20 ATTATTAATTTTAA 7

RESULT 9

AU038857 34 bp mRNA EST 29-MAR-1999
LOCUS AU038857 Dictyostelium discoideum SS (H.Urushihara) Dictyostelium
DEFINITION
discoideum cDNA clone SSL566, mRNA sequence.
ACCESSION AU038857
VERSION AU038857.1 GI:3985610
KEYWORDS EST.
SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum.
REFERENCE Eukaryota; Mycetozoa; Dictyostellida; Dictyostelium.
1 (bases 1 to 34)
Mori, T., Urushihara, H., Salto, T., Ugawa, Y., Mizuno, H., Yoshida, M., Yoshino, R., Mitra, B.N., Pl.M., Sato, T., Takemoto, K., Yasukawa, H., Williams, J., Maeda, M., Takeuchi, T., Ochiai, H. and Tanaka, Y.
The Dictyostelium developmental cDNA project: generation and analysis of expressed sequence tags from the first-finger stage of development
DNA Res. 5 (6), 335-340 (1998)
99156227
Contact: Hideo Urushihara
Institute of Biological Sciences
University of Tsukuba
3-3-10 Ten-nodai, Tsukuba, Ibaraki 305, Japan
Email: d402huesakura.cc.tsukuba.ac.jp
PROJECT = 'Dictyostelium discoideum cDNA project in Japan'.

REFERENCE
AUTHORS Mori, T., Urushihara, H., Salto, T., Ugawa, Y., Mizuno, H., Yoshida, M., Yoshino, R., Mitra, B.N., Pl.M., Sato, T., Takemoto, K., Yasukawa, H., Williams, J., Maeda, M., Takeuchi, T., Ochiai, H. and Tanaka, Y.
The Dictyostelium developmental cDNA project: generation and analysis of expressed sequence tags from the first-finger stage of development
DNA Res. 5 (6), 335-340 (1998)
99156227
Contact: Hideo Urushihara
Institute of Biological Sciences
University of Tsukuba
3-3-10 Ten-nodai, Tsukuba, Ibaraki 305, Japan
Email: d402huesakura.cc.tsukuba.ac.jp
PROJECT = 'Dictyostelium discoideum cDNA project in Japan'.

TITLE

JOURNAL
MEDLINE
COMMENT
Contact: Hideo Urushihara
Institute of Biological Sciences
University of Tsukuba
3-3-10 Ten-nodai, Tsukuba, Ibaraki 305, Japan
Email: d402huesakura.cc.tsukuba.ac.jp
PROJECT = 'Dictyostelium discoideum cDNA project in Japan'.

FEATURES

source

1. .34

Location/Qualifiers

/organism="Dictyostelium discoideum"

/strain="AX4"

/db_xref="taxon:44689"

/clone_lib="SSL566"

/dev_stage="slug"

/dev_stage="slug"

BASE COUNT

23 a 0 c 0 g 11 t

ORIGIN

Query Match
Best Local Similarity 100.0%; Pred. No. 2.5e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 110 ttaataataaaaa 123
|||||
Db 2 TTAATAATAATAAA 15

RESULT 10

AM33885
 LOCUS AM33885 42 bp mRNA EST 31-JAN-2000
 DEFINITION S27B6 AGS-1 Pneumocystis carinii f. sp. carinii cDNA 3', mRNA
 sequence.
 ACCESSION AM33885
 VERSION AM33885.1 GI:6830242
 KEYWORDS EST.
 SOURCE Pneumocystis carinii f. sp. carinii.
 ORGANISM Eukaryota; Fungi; Ascomycota; Pneumocystidomycetes;
 Pneumocystidaceae; Pneumocystis.
 REFERENCE 1 (bases 1 to 42)
 AUTHORS Smilian, A.G., Arnold, J., Weise, M., Wunderlich, J., Staben, C., Edman, J.C., Kovacs, J. and Cushion, M.
 TITLE Expressed sequence tags from Pneumocystis carinii
 JOURNAL Unpublished (2000)
 COMMENT Contact: Staben C
 School of Biological Sciences
 University of Kentucky
 101 Morgan Building, University of Kentucky, Lexington, KY
 40506-0225, USA
 Tel: 606 257 2161
 Fax: 606 257 1717
 Email: staben@pop.uky.edu.
 Location/Qualifiers
 FEATURES
 source 1..42
 /organism="Pneumocystis carinii f. sp. carinii"
 /db_xref="taxon:38081"
 /clone_lib="AGS-1"
 /lab_host="E. coli"
 /note="Vector: lambda ZAP II; Site 1: EcoRI; Site 2: XhoI;
 P. carinii organisms (3x10e9) from a single rat (99-1-6,
 sacrificed on 3/17/99) at Cincinnati VA facilities.
 Triol extracted RNA. Oligo dT priming, standard
 conditions described by vendor, Stratagene. Further
 details see www.uky.edu/Project/Pneumocystis/"

Query Match 4.3%; Score 14; DB 10; Length 42;
 Best Local Similarity 100.0%; Pred. No. 2.4e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 164 ataataaattta 177
 |||||||||||||
 Db 3 ATTAATAAATTTA 16

RESULT 11
 LOCUS AZ634761 42 bp DNA GSS 13-DEC-2000
 DEFINITION IM0490C17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0490C17 R, DNA sequence.
 ACCESSION AZ634761
 VERSION AZ634761.1 GI:11756951
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 42)
 REFERENCE 1 (bases 1 to 42)
 AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
 and Wright, D., Weiss, R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: dunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0490 row: C column: 17
 Seq primer: CACACAGGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 42.
 Location/Qualifiers
 FEATURES
 source 1..42
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone_lib="UUGC1M0490C17"
 /clone_host="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/nares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD2 (g14732114[gb]AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

Query Match 4.3%; Score 14; DB 13; Length 42;
 Best Local Similarity 100.0%; Pred. No. 2.4e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 141 tttttgttttaa 154
 |||||||||||||
 Db 34 TTTTGTGTTTAA 21

RESULT 12
 LOCUS AZ459612 46 bp DNA GSS 04-OCT-2000
 DEFINITION IM0264002R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0264002 R, DNA sequence.
 ACCESSION AZ459612
 VERSION AZ459612.1 GI:10617737
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 46)
 REFERENCE 1 (bases 1 to 46)
 AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
 and Wright, D., Weiss, R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0264 row: 0 column: 02
Seq primer: CACACAGAAACAGCATATGACC
Class: plasmid ends
High quality sequence stop: 46.

FEATURES

source

1. 46
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M0264002"
/clone_1lb="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|9b|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

21 a 5 c 4 g 16 t

ORIGIN

Query Match 4.3%; Score 14; DB 13; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 188 taaataattattt 201

Db 45 TAAATAATTATT 32

RESULT 13

AZ345468 47 bp DNA GSS 29-SEP-2000

LOCUS 1M0080N12F Mouse 10kb plasmid U08C1M library Mus musculus genomic
DEFINITION Clone U08C1M080N12 F, DNA sequence.

ACCESSION AZ345468
VERSION AZ345468.1 GI:10424705

KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 47)
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamli, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss
University of Utah Genome Center

University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0080 row: N column: 12
Seq primer: CGTTGTAACGACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 47.

FEATURES

source

1. 47
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M080N12"
/clone_1lb="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|9b|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

12 a 9 c 0 g 26 t

ORIGIN

Query Match 4.3%; Score 14; DB 13; Length 47;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 112 aataataaaagt 125

Db 26 AATAATAAAAGT 13

RESULT 14

BE043289 50 bp mRNA EST 08-JUN-2000

LOCUS hK49d05.y1 NCI_CGAP_Ov34 Homo sapiens cDNA clone IMAGE:300009 5',
DEFINITION mRNA sequence.

ACCESSION BE043289
VERSION BE043289.1 GI:8360342

KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 50)
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

TITLE Unpublished (1997)

JOURNAL Contact: Robert Strausberg, Ph.D.

COMMENT Email: cgapbs-r@mail.nih.gov
cDNA Library Preparation: David B. Kitzman, Ph.D.
cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL

DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL, send email to:
 info@image.llnl.gov

Seq primer: 40RP from GIBCO.

FEATURES

Location/Qualifiers

```

1..50
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3000009"
/clone_lib="NCL_CGAP_OV34"
/sex="female"
/tissue_type="borderline ovarian carcinoma"
/dev_stage="adult"
/lab_host="DH10B"
/note="Organ: ovary; Vector: pAMP1; mRNA made from
borderline ovarian carcinoma, cDNA made by oligo-OT
priming. Directionally cloned. Size-selected on agarose
gel, average insert size 500 bp. Primary library,
non-amplified."
BASE COUNT      30 a      3 c      11 g      6 t
ORIGIN

```

Query Match 4.3%; Score 14; DB 10; Length 50;
 Best Local Similarity 100.0%; Pred. No. 2.2e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 236 aattgtaaaaaa 249
 |||||
 Db 27 AATTGTAAAAAAA 40

RESULT 15
 LOCUS TA154F10Q 50 bp DNA GSS 13-DEC-2000
 DEFINITION T. brucei sheared genomic DNA clone 154f10, reverse sequence,
 genomic survey sequence.
 ACCESSION AL473287 GI:11838560
 VERSION AL473287.1 GI:11838560
 KEYWORDS GSS.
 SOURCE Trypanosoma brucei.
 ORGANISM Trypanosoma brucei
 Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae;

REFERENCE 1 (bases 1 to 50)
 AUTHORS Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
 Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
 Melville, S.E., Rajadream, M.A. and Barrell, B.G.
 Direct Submission

TITLE Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
 project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
 Cambridgeshire CB10 1SA, E-mail: barrell@sanger.ac.uk and
 nh@sanger.ac.uk

COMMENT Constructed at the Institute for Genomic Research (TIGR),
 Rockville, MD. Genomic DNA isolated from a cloned population of
 Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared
 to give a tight size distribution (4 kb). The v + i method used for the library construction is
 described in detail in Smith, H. and Venter, J.C. (Making small
 insert libraries for whole genome shotgun sequencing projects. In
 Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
 Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org
 Details of T. brucei sequencing at the Sanger Centre are available
 at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

Location/Qualifiers

```

1..50
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="154f10"
BASE COUNT      20 a      7 c      9 g      14 t

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ORIGIN

Query Match 4.3%; Score 14; DB 13; Length 50;
 Best Local Similarity 100.0%; Pred. No. 2.2e+05;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 268 atgaaaaaaatat 281
 |||||
 Db 10 ATGAAAAAATTTAT 23

RESULT 16
 LOCUS A2331628 19 bp DNA GSS 29-SEP-2000
 DEFINITION IM0059M12R Mouse 10kb plasmid UGCM1 library Mus musculus genomic
 clone UGCM1M0059M12 R, DNA sequence.
 ACCESSION A2331628
 VERSION A2331628 GI:10394503
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 19)
 AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
 and Wright, D., Weiss, R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddu@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0059 row: M column: 12
 Seq primer: CACACAGCAACACGATGACC
 Class: plasmid ends
 High quality sequence stop: 19.

FEATURES

Location/Qualifiers

```

1..19
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UGCM1M0059M12"
/clone_lib="Mouse 10kb plasmid UGCM1 library"
/sex="male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F"-
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
digested DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD22 (g114732114[gb]AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

```

```

BASE COUNT      8 a      1 c      0 g      10 t
ORIGIN
Query Match
Best Local Similarity 4.0%; Score 13; DB 13; Length 19;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 166 aaataaattag 178
Db 14 AAATAAATTAG 2

RESULT 17
AZ829725 25 bp DNA GSS 20-FEB-2001
LOCUS 2M0107124F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC2M0107124 F, DNA sequence.
ACCESSION AZ829725
VERSION AZ829725.1 GI:1299549
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 25)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0107 row: I column: 24
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers
1..25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0107124"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapored DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (9114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapored mouse DNA was annealed to
adapored vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells

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```

BASE COUNT      14 a      3 c      4 g      4 t
ORIGIN
Query Match
Best Local Similarity 4.0%; Score 13; DB 13; Length 25;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 301 ttatgtttcat 313
Db 24 TTTATGTTTCAT 12

RESULT 18
AZ309204 26 bp DNA GSS 29-SEP-2000
LOCUS 1M0013F07F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC1M0013F07 F, DNA sequence.
ACCESSION AZ309204
VERSION AZ309204.1 GI:10349955
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 26)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0013 row: F column: 07
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 26.
Location/Qualifiers
1..26
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0013F07"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapored DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (9114732114|9b|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapored mouse DNA was annealed to
adapored vector DNA, and transformed into

```

chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208

Db 7 TTATTTTATTTT 19

RESULT 19
LOCUS A2866662/c 26 bp DNA GSS 21-FEB-2001
DEFINITION 2M0177A18F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0177A18 F, DNA sequence.

ACCESSION A2866662

VERSION A2866662.1 GI:13068193

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 26)

1 (bases 1 to 26)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0177 row: A column: 18

Seq primer: CGTGTAAACGACGCCACG

Class: plasmid ends

High quality sequence stop: 26.

Location/Qualifiers

1..26

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC2M0177A18"

/clone_id="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g114732114gblAF129072.1), a copy-number
inducible derivative of plasmid RI. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttt 208

Db 17 TTATTTTATTTT 5

RESULT 20
LOCUS TA123B120/c 26 bp DNA GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 123b12, reverse sequence,
genomic survey sequence.

ACCESSION AL463522

VERSION AL463522.1 GI:11834032

KEYWORDS GSS.

SOURCE Trypanosoma brucei.

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.

1 (bases 1 to 26)

Hall,N., Bowman,S., Leonard,N.J., Doggett,J., Atkin,R.,
Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,
Melville,S.E., Rajandream,M.A. and Barrell,B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
nh@sanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 G9nat 10.1) was mechanically sheared
to give a tight size distribution (4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaubin and B.
Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org

Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/Projects/T_brucei/.

Location/Qualifiers

1..26

/organism="Trypanosoma brucei"

/strain="TREU927"

/db_xref="taxon:5691"

/clone="123b12"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g114732114gblAF129072.1), a copy-number
inducible derivative of plasmid RI. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to

Query Match 4.0%; Score 13; DB 13; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 271 aaaaaattattt 283

Db 20 AAAAAATTATTT 8

RESULT 21
LOCUS A2784620 27 bp DNA GSS 16-FEB-2001
DEFINITION 2M0027P04R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0027P04 R, DNA sequence.

ACCESSION A2784620

VERSION A2784620.1 GI:11834032

KEYWORDS GSS.

SOURCE Trypanosoma brucei.

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.

1 (bases 1 to 26)

Hall,N., Bowman,S., Leonard,N.J., Doggett,J., Atkin,R.,
Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,
Melville,S.E., Rajandream,M.A. and Barrell,B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
nh@sanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 G9nat 10.1) was mechanically sheared
to give a tight size distribution (4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaubin and B.
Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org

Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/Projects/T_brucei/.

Location/Qualifiers

1..26

/organism="Trypanosoma brucei"

/strain="TREU927"

/db_xref="taxon:5691"

/clone="123b12"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"

/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g114732114gblAF129072.1), a copy-number
inducible derivative of plasmid RI. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to

ACCESSION A2784620
 VERSION A2784620.1 GI:12920544
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamli,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0027 row: P column: 04
 Seq primer: CACACAGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 27.
 FEATURES
 source
 1..27
 Location/Qualifiers
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M0027P04"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
 BASE COUNT
 5 a 5 c 5 g 12 t
 ORIGIN
 Query Match 4.0%; Score 13; DB 13; Length 27;
 Best Local Similarity 100.0%; Pred. No. 6.8e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 137 aaacttttttgt 149
 ||||||||||||
 Db 15 ACACTTTTGT 27
 RESULT 22
 A2452653 28 bp DNA GSS 04-OCT-2000
 LOCUS A2452653
 DEFINITION 1M0252E07R Mouse 10kb plasmid UUGC1M library Mus musculus genomic

clone UUGC1M0252E07 R, DNA sequence.
 ACCESSION A2452653
 VERSION A2452653.1 GI:10609676
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamli,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0252 row: E column: 07
 Seq primer: CACACAGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 28.
 FEATURES
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 1..28
 Location/Qualifiers
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0252E07"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
 BASE COUNT
 14 a 2 c 2 g 10 t
 ORIGIN
 Query Match 4.0%; Score 13; DB 13; Length 28;
 Best Local Similarity 100.0%; Pred. No. 6.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 271 aaaaaaatattt 283
 ||||||||||||
 Db 16 AAAAAATATT 28
 RESULT 23
 A2623794 30 bp DNA GSS 13-DEC-2000
 LOCUS A2623794/C

DEFINITION 1M0461C19R Mouse 10kb plasmid U06C1M library Mus musculus genomic clone U06C1M0461C19 R, DNA sequence.

ACCESSION AZ623794

VERSION AZ623794.1 GI:11745984

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 30)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamli, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A. and Wright, D., Weiss, R.

AUTHORS Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

TITLE

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss
University of Utah
Department of Molecular Biology
Room 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0461 row: C column: 19
Seq primer: CACACAGGAAACGCTATGACC
Class: plasmid ends
High quality sequence stop: 30.
Location/Qualifiers

FEATURES

SOURCE

1. 30
location/Qualifiers

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C1M0461C19"
/clone_1lb="Mouse 10kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b) (AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 4 c 3 g 9 t

ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.5e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 202 ttatttaattt 214
|||||

Db 14 TTATTTAAATTT 2

RESULT 24
H40874

LOCUS H40874 32 bp mRNA EST 31-JUL-1995

DEFINITION yp97e10.s1 Soares adult brain N2b5HB55y Homo sapiens cDNA clone IMAGE:176394.3, similar to gb:X54150.rna1 1M0N0GLOBULIN ALPHA FC RECEPTOR PRECURSOR (HUMAN);, mRNA sequence.

ACCESSION H40874

VERSION H40874.1 GI:916926

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 32)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kneaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevisakis, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.

AUTHORS The Washu-Merck EST Project

TITLE

JOURNAL Unpublished (1995)

COMMENT Contact: Wilson R.
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert size: 674
High quality sequence starts: 1
High quality sequence stops: 1
Source: IMAGE Consortium, LNL
This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert length: 674 Std Error: 0.00
Seq primer: Promega -21m13
High quality sequence stop: 1.
Location/Qualifiers

FEATURES

SOURCE

1. 32
location/Qualifiers

/organism="Homo sapiens"
/db_xref="GDB:3838590"
/db_xref="taxon:9606"
/clone="IMAGE:176394"
/clone_1lb="Soares adult brain N2b5HB55y"
/sex="Male"
/dev_stage="55-year old"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: brain; Vector: pT73D (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCTGAAGTGGAGCGCGCGCTTTTATTTTATTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library went through one round of normalization to a Cot = 53. Library constructed by Bento Soares and M. Fatima Bonaldo. The adult brain RNA was provided by Dr. Donald H. Gilden. Tissue was acquired 17-18 hours after death which occurred in consequence of a ruptured aortic aneurysm. RNA was prepared from a pool of tissues representing the following areas of the brain: frontal, parietal, temporal and occipital cortex from the left and right hemispheres, subcortical white matter, basal ganglia, thalamus, cerebellum, midbrain, pons and medulla."

BASE COUNT 6 a 0 c 0 g 26 t

ORIGIN

Query Match 4.0%; Score 13; DB 11; Length 32;
Best Local Similarity 100.0%; Pred. No. 6.4e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttattt 208
|||||

Db 15 TTATTTTATTTT 27

```

RESULT 25
LOCUS      AZ458690
DEFINITION 1M0263E04F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
ACCESSION  AZ458690
VERSION     AZ458690.1
KEYWORDS    GI:10616815
SOURCE      house mouse.
ORGANISM    Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE   1 (bases 1 to 32)
AUTHORS     Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
            ,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
            and Wright,D., Weiss,R.
TITLE       Mouse whole genome scaffolding with paired end reads from 10kb
            plasmid inserts
JOURNAL     Unpublished (2000)
COMMENT     Contact: Robert B. Weiss
            University of Utah Genome Center
            University of Utah
            Rm. 306, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
            84112, USA
            Tel: 801 585 5606
            Fax: 801 585 7177
            Email: ddunn@genetics.utah.edu
            Insert Length: 10000 Std Error: 0.00
            Plate: 0263 row: E column: 04
            Seq primer: CGTGTGAAACGACGCGCCAGT
            Class: plasmid ends
            High quality sequence stop: 32.
FEATURES
source
1..32
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_1ib="UUGC1M0263E04"
/clone_1ib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42ny; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (q11473214|9b|AF129072.1), a copy-number
inducible derivative of plasmid RL. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
BASE COUNT      4 a
ORIGIN          5 c
                4 g
                19 t
Query Match
Best Local Similarity 100.0%; Score 13; DB 13; Length 32;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
196 ttaatttttttt 208
|||||

```

```

Db 12 TTATTTTATTTT 24
RESULT 26
LOCUS      AA906810/C
DEFINITION AA906810 34 bp mRNA
ACCESSION  AA906810
VERSION     AA906810.1
KEYWORDS    GI:3042054
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE   1 (bases 1 to 34)
AUTHORS     NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE       National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL     Unpublished (1997)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
            Emmert-Buck, M.D., Ph.D.
            CDNA Library Preparation by: Bento Soares, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Trace considered overall poor quality
            Seq primer: -40m13 fwd. ET from Amerisham
            High quality sequence stop: 1.
FEATURES
source
1..34
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_1ib="IMAGE:1519586"
/clone_1ib="NCI-CGAP-GC4"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker. 1st strand cDNA was prepared from 3 pooled
germ cell tumors, and was then primed with a Not I -
oligo(dT) primer. Double-stranded cDNA was ligated to Eco
RI adaptors (Pharmacia), digested with Not I and cloned
into the Not I and Eco RI sites of the modified pT7T3
vector. Library is normalized. Library was constructed by
Bento Soares and M. Fatima Bonaldo."
BASE COUNT      8 a
ORIGIN          4 c
                7 g
                15 t
Query Match
Best Local Similarity 100.0%; Score 13; DB 10; Length 34;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
51 ataataaagaat 63
|||||
Db 21 ATATATATAAAGAT 9
RESULT 27
LOCUS      AZ586746
DEFINITION 1M0392K22R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
ACCESSION  AZ586746
VERSION     AZ586746.1
KEYWORDS    GI:11708936
SOURCE      house mouse.
Query Match
Best Local Similarity 100.0%; Score 13; DB 13; Length 32;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
196 ttaatttttttt 208
|||||

```


ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 34)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0392 row: K column: 22
Seq primer: CACACAGCAACACGTATGACC
Class: plasmid ends
High quality sequence stop: 34.
Location/Qualifiers
1..34
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C1M0392K22"
/clone_lib="Mouse 10kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gblAF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 2 a 0 c 4 g 28 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 34;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttatttt 208
|||||
Db 6 TTATTTTATTTT 18

RESULT 28
A2781725 34 bp DNA GSS 16-FEB-2001
LOCUS
DEFINITION 2M002111AF Mouse 10kb plasmid U06C1M library Mus musculus genomic
ACCESSION A2781725
VERSION A2781725.1 GI:12914706
KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 34)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0021 row: I column: 14
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 34.
Location/Qualifiers
1..34
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C2M0021114"
/clone_lib="Mouse 10kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gblAF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 13 a 4 c 4 g 13 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 34;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 209 aaattgttaaaa 221
|||||
Db 10 AAATTGTTAAAA 22

RESULT 29
AM246486 35 bp mRNA EST 07-JAN-2000
LOCUS
DEFINITION 2821345.3pr1me NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2821345 3',
ACCESSION AM246486
VERSION AM246486.1 GI:6589479

KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
TITLE 1 (bases 1 to 35)
JOURNAL NIH-MGC http://mgc.nci.nih.gov/.
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Other_ESTS: 2821545.5prime
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: DCTD/DPB CDNA Library Preparation; Lung
Hong/Rubin Laboratory CDNA Library Arrayed by: The I.M.A.G.E.
Consortium (LNL) DNA sequencing by: Berkeley MGC sequencing
project clone distribution: MGC clone distribution information can
be found through the I.M.A.G.E. Consortium/LNL at:
www.bio.lnl.gov/dbp/image/image.html Base Calling / Quality
Scores: PHRED from University of Washington Genome Center. Vector
Trimming: crossmatch from University of Washington genome center
PHRAP suite. Poly-T identification: patmatch.pl from Berkeley
Drosophila Genome Project. University of Washington Genome Center:
http://www.genome.washington.edu Low Quality Sequence: 35
contiguous PHRED high quality bases following vector sequence. Very
low quality sequence: Trace file contained 35 contiguous distinct
peaks following vector sequence. Polyadenylation: Based upon the
presence of a XhoI site followed by a run of 14 or more T residues
at the beginning of the sequence, this CDNA insert was
polyadenylated.
Plate: LCM7 row: B column: 10
High quality sequence stop: 35.
Location/Qualifiers
1..35
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="NIH-MGC-7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/note="Organ: Lung; Vector: pOT7; Site_1: XhoI; Site_2:
EcoRI; CDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 11 a 4 c 1 g 19 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 35;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 35 tttaaatgaata 47
|||||
Db 8 TTTAAATGAATA 20

RESULT 30
LOCUS D45807 35 bp mRNA EST 20-FEB-1995
DEFINITION HKMS0025 Human adult lung 3' directed MboI CDNA Homo sapiens CDNA
3', mRNA sequence.
ACCESSION D45807
VERSION D45807.1 GI:662761
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 35)
AUTHORS Itoh, K., Okubo, K., Yosi, J., Yokouchi, H. and Matsubara, K.
TITLE An expression profile of active genes in human lung
JOURNAL DNA Research 1, 279-287 (1994)
MEDLINE 95236275
COMMENT Contact: Kohichi Itoh
Institute for Molecular and Cellular Biology
Osaka University
3-1, Yamadaoka, Suita, Osaka, 565, Japan
Tel: 06-877-5111 x3910
Fax: 06-877-1922.
FEATURES
source
1..35
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Human adult lung 3' directed MboI CDNA"
/note="Adult human lung, 3' directed MboI"
BASE COUNT 14 a 3 c 5 g 13 t
ORIGIN
Query Match 4.0%; Score 13; DB 11; Length 35;
Best Local Similarity 100.0%; Pred. No. 6.2e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 106 atagtaataaat 118
|||||
Db 11 ATAGTTAATAAT 23

RESULT 31
LOCUS AZ314238 36 bp DNA GSS 29-SEP-2000
DEFINITION 1M0030N24R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
clone UUCG1M0030N24 R, DNA sequence.
ACCESSION AZ314238
VERSION AZ314238.1 GI:10359929
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 36)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamli, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Rilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10Kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0030 row: N column: 24
Seq primer: CACACGAGAACACGCTATGACC
Class: plasmid ends
High quality sequence stop: 36.
Location/Qualifiers
1..36
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_lib="UUCG1M0030N24"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g14732114[gb|AF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 8 a 3 c 5 g 20 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 36;
Best Local Similarity 100.0%; Pred. No. 6;le+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 312 atgtttcttatt 324
|||||
Db 2 ATGTTTCTATT 14

RESULT 32
LOCUS AA913140 37 bp mRNA 14-APR-1998
DEFINITION o118h06.s1 NCI-CGAP_HNI Homo sapiens cDNA IMAGE:1483067 3'
similar to SW:YK13 YEAST P36079 HYPOTHETICAL 23.7 KD PROTEIN IN
MDH1-VMA5 INTERGENIC REGION. ;, mRNA sequence.
AA913140
ACCESSION AA913140.1 GI:3052532
VERSION EST.
KEYWORDS human.
SOURCE Human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 37)
NCI/NIH-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute / National Institute of Dental Research,
Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
Unpublished (1997)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgapbs-rt@mail.nih.gov
Tissue Procurement: John Ensley, M.D., Mary May, J. Silvio Gutkind, Ph.D.

CDNA Library Preparation: Stratagene, Inc.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:
www-bio.llnl.gov/bdrip/image/image.html

Trace considered overall poor quality
Seq primer: -40m13 fwd. RT from Amersham
High quality sequence stop: 1.

FEATURES

Source
1. .37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1483067"
/clone_lib="NCI-CGAP_HNI"
/tissue_type="squamous cell carcinoma"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: lymph node; Vector: Bluescript SK-; Site_1: EcoRI; Site_2: XhoI; Cloned unidirectionally. Primer:

OLigo dT. Average insert size 1.3 kb. 5' adaptor sequence:
5' GAATTCGGCAGCAG 3' 3' adaptor sequence: 5' (GA
1)ACATGATCTCGAGTTTCTTTTCTTTT 3' "

BASE COUNT 2 a 9 c 2 g 24 t
ORIGIN

Query Match 4.0%; Score 13; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttatttt 208
|||||
Db 3 TTATTTTATT 15

RESULT 33
LOCUS AU009123 37 bp mRNA 31-JUL-1998
DEFINITION AU009123 Schizosaccharomyces pombe late log phase cDNA
Schizosaccharomyces pombe cDNA spec04510, mRNA sequence.
AU009123
ACCESSION AU009123 GI:3345803
VERSION EST.
KEYWORDS fission yeast.
SOURCE Schizosaccharomyces pombe
ORGANISM Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
Schizosaccharomycetaceae; Schizosaccharomycetaceae;
Schizosaccharomycetes.
1 (bases 1 to 37)
REFERENCE Moriwo,M. and Mita,K.
AUTHORS Identification of expressed sequence tags of Schizosaccharomyces
TITLE Pombe
JOURNAL Unpublished (1998)
COMMENT Contact: MitsuoKI Moriwo
Genome Research Group
National Institute of Radiological Sciences
9-1, Anagawa 4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: moriwo@nirs.go.jp.
Location/Qualifiers

FEATURES

source
1. .37
/organism="Schizosaccharomyces pombe"
/strain="972"
/db_xref="taxon:4896"
/clone="spc04510"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/sex="h minus"
/note="Vector: M13mp19; The cDNA library of
Schizosaccharomyces pombe was prepared by cloning cDNA
into the SmaI site of M13mp19 DNA and the direction of DNA
sequences was not always from 5' to 3'. The cDNA data of
Schizosaccharomyces pombe are available for searching on
the World Wide Web. (URL, <http://www.nirs.go.jp>)"
BASE COUNT 13 a 6 c 3 g 15 t
ORIGIN

Query Match 4.0%; Score 13; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 110 ttataataaaaa 122
|||||
Db 13 TTAATAATAAAA 25

RESULT 34
LOCUS AA916625 40 bp mRNA 10-JUN-1998
DEFINITION om05g12.s1 Soares_MFL_T_GRC.S1 Homo sapiens cDNA clone
IMAGE:1540198 3 similar to SW:NO5M.TRYBB P04540 NADH-UBIQUINONE
OXIDOREDUCTASE CHAIN 5 ;, mRNA sequence.
AA916625

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VERSION AA916625.1 GI:3056017
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 40)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 1026 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 1.
FEATURES
source
1..40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1540198"
/clone_1lb="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site: 1: Not I; Site: 2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBH139, testis NHT, and B-cell
NCI-CGAP-CGI) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
1.M.A.G.E. clones 297480-302087, 662632-667239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo. "
BASE COUNT 10 a 1 c 2 g 27 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 40;
Best Local Similarity 100.0%; Pred. No. 5.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 106 atagtgtaataat 118
|||||
Db 21 ATAGTAAATAAAT 9

RESULT 35
LOCUS AA922076 40 bp mRNA EST 21-APR-1998
DEFINITION Oh08907.s1 NCI-CGAP Co8 Homo sapiens cDNA clone IMAGE:1457244 3'
similar to TR:Q34096 Q34096 WURF2 PROTEIN. ;, human sequence.
ACCESSION AA922076
VERSION AA922076.1 GI:3069385
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 40)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.

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cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/dbfp/image/image.html
Trace considered overall poor quality
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 1.
FEATURES
source
1..40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1457244"
/clone_1lb="NCI-CGAP Co8"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"
/note="Organ: colon; Vector: pT7T3D-Pac (Pharmacia) with a
modified polylinker; 1st strand cDNA was prepared from
colon adenocarcinoma, and was then primed with a Not I -
oligo(drf) primer. Double-stranded cDNA was ligated to Eco
RI adaptors (Pharmacia), digested with Not I and cloned
into the Not I and Eco RI sites of the modified pT7T3
vector. Library is normalized. Library was constructed by
Bento Soares and M. Fatima Bonaldo. "
BASE COUNT 5 a 4 c 3 g 28 t
ORIGIN
Query Match 4.0%; Score 13; DB 10; Length 40;
Best Local Similarity 100.0%; Pred. No. 5.9e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 112 aataataaaag 124
|||||
Db 38 AATTAATAAAG 26

RESULT 36
LOCUS C00434 41 bp mRNA EST 23-JUL-1996
DEFINITION HUMGS0006099 Human adult (K.Okubo) Homo sapiens cDNA, mRNA
sequence.
ACCESSION C00434
VERSION C00434.1 GI:1432664
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 41)
AUTHORS Okubo, K.
TITLE BodyMap: human gene expression database
JOURNAL Unpublished (1995)
COMMENT Contact: Okubo, K.
Institute for Molecular and Cellular Biol
Osaka University
1-3 Yamada-oka, Suita, Osaka Pref. 565, Japan
Tel: 06-877-5111(ex.3315)
Email: kousaku@imcb.osaka-u.ac.jp
Human Gene Signature, 3'-directed cDNA sequence. We are not
submitting the same cDNA sequence redundantly to DDBJ since 1993.
For the abundance information of clones with this sequence in this
library and as well as in other 3'-directed libraries, see
http://www.imcb.osaka-u.ac.jp/bodymap/. The sequences of the clones
represented by this GS sequences is also found there.
FEATURES
source
1..41
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_1lb="Human adult (K.Okubo)"
/dev_stage="adult"
BASE COUNT 12 a 5 c 7 g 17 t

```

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ORIGIN

Query Match      4.0%; Score 13; DB 11; Length 41;
Best Local Similarity 100.0%; Pred. No. 5.8e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 109 gttataataaa 121
Db 14 GTTAATAATAA 26

RESULT 37
A2777050/c 41 bp DNA 16-FEB-2001
LOCUS 2M001M12F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC2M001M12 F, DNA sequence.
ACCESSION A2777050
VERSION A2777050.1 GI:12905260
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 41)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLc, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0011 row: M column: 12
Seq primer: CGTGTAAACGACGCCACGT
Class: plasmid ends
High quality sequence stop: 41.
Location/Qualifiers
1. 41
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M001M12"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g14732114[gbl]AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

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BASE COUNT      9 a      3 c      12 g      17 t
ORIGIN

Query Match      4.0%; Score 13; DB 13; Length 41;
Best Local Similarity 100.0%; Pred. No. 5.8e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 57 aaaagatatata 69
Db 16 AAAAGATATATTA 4

RESULT 38
A2371136 43 bp DNA 02-OCT-2000
LOCUS 1M0122N05F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION clone UUGC1M0122N05 F, DNA sequence.
ACCESSION A2371136
VERSION A2371136.1 GI:10484836
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 43)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLc, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0122 row: N column: 05
Seq primer: CGTGTAAACGACGCCACGT
Class: plasmid ends
High quality sequence stop: 43.
Location/Qualifiers
1. 43
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0122N05"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PMD42 (g14732114[gbl]AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells

```

BASE COUNT 28 a 2 c 2 g 11 t
 ORIGIN and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 111 taataataaaaa 123
 Db 18 TAATAAATAAAAA 30

RESULT 39
 AZ464392 43 bp DNA GSS 04-OCT-2000
 LOCUS 1M0273D17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 DEFINITION clone UUGC1M0273D17 R, DNA sequence.
 ACCESSION AZ464392
 VERSION AZ464392.1 GI:10622517
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 43)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Haml, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A.
 and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0273 row: D column: 17
 Seq primer: CACACAGAAACAGCATGACC
 Class: plasmid ends
 High quality sequence stop: 43.
 FEATURES
 source Location/Qualifiers
 1..43
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0273D17"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g114732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into

BASE COUNT 6 a 7 c 8 g 22 t
 ORIGIN chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 139 acctttttgtttt 151
 Db 30 ACTTTTGTGTTTT 42

RESULT 40
 AZ575514 43 bp DNA GSS 06-DEC-2000
 LOCUS AST-T21F0045 Genetrap T47D Human Breast Carcinoma Library Homo
 DEFINITION sapiens genomic 5', DNA sequence.
 ACCESSION AZ575514
 VERSION AZ575514.1 GI:11561825
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 1 (bases 1 to 43)
 Henkel, G., Liyanage, M., Pratt, E., Huang, D., Riley, M., Bernardino, A.,
 Durick, K., and Pollok, B.
 Exon-trap tags from a T47D Genomescreen(TM) Library
 Unpublished (2000)
 JOURNAL Contact: Greg Henkel
 COMMENT Gene Expression
 Aurora Biosciences Corp.
 11010 Torreyana Road, San Diego, CA 92121, USA
 Tel: 8584048436
 Fax: 8584046719
 Email: henkelg@aurorabio.com
 Pools of cells were isolated from a GenomesScreen(TM) library. The
 library of cells was generated by retroviral integration of a gene
 tagging element consisting of: 1) A promoterless beta-lactamase
 proceeded by a splice acceptor as a reporter for gene expression;
 2) A promoter driving neomycin resistance followed by a splice
 donor to trap downstream exons. 3' RACE from neomycin gene was
 performed using total RNA from isolated pools. Output was shotgun
 cloned in pAMP-1 and used to transform DH5-alpha competent
 bacteria. 5' ends of reported sequences were immediately preceded
 by splice donor from the trapping construct.
 Class: exon-trapped.
 FEATURES
 source Location/Qualifiers
 1..43
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="Genetrap T47D Human Breast Carcinoma Library"
 /tissue_type="Carcinoma"
 /cell_type="Epithelial"
 /cell_line="T47D"
 /note="Organ: Breast; Vector: pAMP-1; 3' RACE of total RNA
 from genetrap pools; shotgun clone in pAMP-1 and used to
 transform DH5-alpha competent bacteria."

BASE COUNT 18 a 9 c 5 g 11 t

Query Match 4.0%; Score 13; DB 13; Length 43;
 Best Local Similarity 100.0%; Pred. No. 5.7e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 39 aatgaataaaga 51
 Db 1 AATGAATAAAGA 13

RESULT 41
D20668/c 45 bp mRNA EST 30-JUL-1996
DEFINITION HM06501644 Human promyelocyte Homo sapiens cDNA clone pm2268 3',
mRNA sequence.
ACCESSION D20668
VERSION D20668.1 GI:501764
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS Okubo, K., Fukushima, A., Yoshi, J., Niiyama, T., Kojima, Y., Yoshinari,
H., Arimoto, J., and Matsubara, K.
TITLE Gene expression of human promyelocytic cell line HL60 before and
after induction of differentiation. A new application of 3'directed
cDNA sequencing
Unpublished (1993)
JOURNAL
COMMENT Contact: Okubo, K., Fukushima, A., Yoshi, J., Niiyama, T., Kojima, Y.,
Yoshinari, H., Arimoto, J., and Matsubara, K.
Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
FEATURES
SOURCE Location/Qualifiers
1..45
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="pm2268"
/clone_lib="Human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type =
promyelocyte."
BASE COUNT 17 a 6 c 6 g 14 t 2 others
ORIGIN

Query Match 4.0%; Score 13; DB 11; Length 45;
Best Local Similarity 100.0%; Pred. No. 5.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 196 ttattttatttt 208
|||||
Db 34 TTATTTTATTTT 22

RESULT 42
A2459612 46 bp DNA GSS 04-OCT-2000
LOCUS A2459612
DEFINITION HM0264002R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0264002 R, DNA sequence.
ACCESSION A2459612
VERSION A2459612.1 GI:10617737
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Rellly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunne@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0264 row: 0 column: 02
Seq primer: CACACAGCAACACGATGACC
Class: plasmid ends
High quality sequence stop: 46.
Location/Qualifiers
1..46
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0264002"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptored DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (91473211419b/AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptored mouse DNA was annealed to
adaptored vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 21 a 5 c 4 g 16 t
ORIGIN

Query Match 4.0%; Score 13; DB 13; Length 46;
Best Local Similarity 100.0%; Pred. No. 5.6e+05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 190 aaataattattt 202
|||||
Db 32 AAATAATTATTTT 44

RESULT 43
A2834972 46 bp DNA GSS 20-FEB-2001
LOCUS A2834972/c
DEFINITION 2M0117N19R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0117N19 R, DNA sequence.
ACCESSION A2834972
VERSION A2834972.1 GI:13004880
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Rellly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
JOURNAL
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606

Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0117 row: N column: 19
 Seq primer: CACACAGCAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 46.

FEATURES

SOURCE

Location/Qualifiers

1. 46
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M0117N19"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

15 a 7 c 9 g 15 t

BASE COUNT

23 a 10 c 5 g 8 t

Query Match
 Best Local Similarity 100.0%; Score 13; DB 13; Length 46;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 30 gaatgttaaatc 42
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 Db 20 GAATGTTTAAATG 8

RESULT 44

AZ991460 46 bp DNA GSS 27-APR-2001
 DEFINITION 2M0275C07R Mouse 10kb plasmid UUGC2M library Mus musculus genomic
 accession AZ991460
 version AZ991460.1 GI:13862687
 keywords GSS.
 source house mouse.
 organism Mus musculus

REFERENCE
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 unpublished (2000)

TITLE

JOURNAL

COMMENT

Contact: Robert B. Weiss
 University of Utah Genome Center
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA

Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0275 row: K column: 07
 Seq primer: CACACAGCAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 46.

FEATURES

SOURCE

Location/Qualifiers

1. 46
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M0275K07"
 /clone_lib="Mouse 10kb plasmid UUGC2M library"
 /sex="Female"
 /lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT

23 a 10 c 5 g 8 t

Query Match
 Best Local Similarity 100.0%; Score 13; DB 13; Length 46;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 taatgaataaa 49
 |||||
 Db 32 TAAATGAATAAAA 44

RESULT 45

A1630064 49 bp mRNA EST 08-MAR-2000
 DEFINITION ad00157 Proliferating Erythroid cells (LCB-ad library) Homo sapiens
 accession A1630064
 version A1630064.1 GI:4681394
 keywords EST.
 source human.
 organism Homo sapiens

REFERENCE
 AUTHORS Gubin,A.N., Miyoro,J.M., Bouffard,G.G. and Miller,J.L.
 Gene expression in proliferating human erythroid cells
 Genomics 59 (2), 168-177 (1999)

TITLE

JOURNAL

COMMENT

Contact: Jeffery L. Miller
 Laboratory of Chemical Biology
 National Institute of Diabetes and Digestive and Kidney Diseases
 Building 10, Room 9B17, National Institutes of Health, Bethesda, MD 20892, USA
 Tel: 301 402 2373
 Fax: 301 435 5148

Email: jmf@nih.gov
 The 'ad' library was constructed by Alexander Gubin, Ph.D. in the
 Laboratory of Chemical Biology, NIDDK, NIH. DNA Sequencing and/or
 analyses by National Institutes of Health Intramural Sequencing
 Center (NISC). More information available at:
<http://hembase.nidck.nih.gov>.

FEATURES

source

```

1. .49
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="ad00157"
/clone_lib="Proliferating Erythroid Cells (LCB:ad library)"
/sex="unknown"
/tissue_type="blood"
/cell_type="Erythroid Cells"
/cell_line="Primary Culture of Peripheral Blood Mononuclear Cells"
/dev_stage="Progenitor; EPO responsive CD71+++"
/lab_host="DH5alpha"
/note="Organ: blood; Vector: pCRIT; Site_1: EcORI; Site_2: EcORI; Human peripheral blood mononuclear cells were cultured in the presence or absence of erythropoietin. CD71+++ cells arising only in erythropoietin-supplemented medium were isolated by fluorescence activated cell sorting. Those cells demonstrated an average of six additional doublings in suspension culture and erythroid colony formation in methylcellulose. Suppression subtractive hybridization was used to construct the ad library (tester-sorted CD71+++ cells, driver=unsorted cells cultured without erythropoietin)."
BASE COUNT      16 a      7 c      9 g     13 t      4 others
ORIGIN

```

Query Match 4.0%; Score 13; DB 10; Length 49;
 Best Local Similarity 100.0%; Pred. No. 5.4e+05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 172 aattagataaaa 184
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 Db 24 AATTGATGATFAAAA 36

Search completed: January 24, 2002, 02:57:29
 Job time: 3951 sec

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[illegible]

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Best Local Similarity	49.7%	Pred. NO. 0.021;		
Matches 151; Conservative	0;	Mismatches 153;	Indels 0;	Gaps
0y	24	acagaagaatgtttaatgaataaagaataaanaagaataataataataagctgaa	83	

Db 1878 AAAAAAAAAATTTTAAATTTAAAAAGAAAAAGAAAAAGTACGATTATTA 1819
QY 84 aattatattatagatagatagatagatagatagatagatagatagatagat 143
Db 1818 AAAAAATTAATATTTTTCATCTTAATTAATTAATTAATTAATTAATTAAT 1759
QY 144 ttgttttaaaaggaaataataaattagataaaagtgtaaaataattttt 203
Db 1758 TATATCTATCTAAAAAAGAGGAAAGATGATGATGATGATGATGATGATGAT 1699
QY 204 atttaaatctgttaaaatttgatataattgaaattgttaaaaaaattcagg 263
Db 1698 TTTTATTTTATTTTAAAAAGAGTCTCATGACAAAAAGTGTCTCATGACAAA 1639
QY 264 ataaatgaaaaaattatttcaagtttactgttaatttttattgttttcatgt 323
Db 1638 AAAAAAAGAGGAGGAAAGTAAATTAATTAATTAATTAATTAATTAATTAAT 1579
QY 324 tgtt 327
Db 1578 ATTT 1575

RESULT 6
US-08-973-273-4/c
Sequence 4, Application US/08973273
Patent No. 6140085
GENERAL INFORMATION:
APPLICANT: Dean, Caroline
APPLICANT: MacKnight, Richard C
APPLICANT: Bancroft, Ian
APPLICANT: Lister, Clare K
TITLE OF INVENTION: Genetic Control of Flowering
NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESSEE: Nixon & Vanderyhe P.C.
STREET: 1100 No. 6140085th Glebe Road, 8th Floor
CITY: Arlington
STATE: Virginia
COUNTRY: United States of America
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/973,273
FILING DATE: 01-DEC-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/GB96/01332
FILING DATE: 03-JUN-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 951196.9
FILING DATE: 02-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: Ms Mary J Wilson
REGISTRATION NUMBER: 32,955
REFERENCE/DOCKET NUMBER: 620-29
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 816-4000
TELEFAX: (703) 816-4100
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 9048 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Brassica

US-08-973-273-4
Query Match 17.8%; Score 58.2; DB 3; Length 9048;
Best Local Similarity 53.5%; Pred. No. 0.029;
Matches 144; Conservative 0; Mismatches 123; Indels 2; Gaps 1;
QY 35 tttaataaataaagaataaataaataaataaataaataaataaataaataa 94
Db 452 TCTTATATGATTTATATATATTTAAACGATTTTATTTATATCAGGAAATATATATGTTA 393
QY 95 atatgataagtagtataataaagaatgttctcgggggacacttttgttttaa 154
Db 392 TATTAATATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 333
QY 155 aaggaataataaataaataaattagataaaagtgttaaaataatttttattt 214
Db 332 AAGTTAATTTCTATATATATTTTGTGCTATCTGAAACAA--TATTTTATATTAATAAGT 275
QY 215 gttaaaatttgatataattgtaattgttaaaaaaatttcagggggaatataa 274
Db 274 TAAAAACATTTATATTAAGATATTTTCTTAACCTATTTCTAGATATGAGTGTTTAA 215
QY 275 aaattattcaagtttactgttaatttt 303
Db 214 AATTTAACACATTAATTAATTAATTTT 186

RESULT 7
US-08-998-416-1137/c
Sequence 1137, Application US/08998416
Patent No. 6239264
GENERAL INFORMATION:
APPLICANT: Philippsen, Peter
APPLICANT: Pohlmann, Rainer
APPLICANT: Steiner, Sabine
APPLICANT: Mohr, Christine
APPLICANT: Wendland, Jurgen
APPLICANT: Knechtle, Philipp
APPLICANT: Redischung, Corinne
TITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSYPPII
NUMBER OF SEQUENCES: 1152
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 6239264rtis Corporation
STREET: 3054 Cornwalis Road
CITY: Research Triangle Park
STATE: No. 6239264th Carolina
COUNTRY: USA
ZIP: 27709
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: CH 0016/97
FILING DATE: 31-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Meigs, J. Timothy
REGISTRATION NUMBER: 38,241
REFERENCE/DOCKET NUMBER: PF/5-30306/A/CGC1976
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8689
TELEFAX: 919-541-8689
INFORMATION FOR SEQ ID NO: 1137:
SEQUENCE CHARACTERISTICS:
LENGTH: 636 base pairs
TYPE: nucleic acid

GENERAL INFORMATION:
APPLICANT: Philippsen, Peter
APPLICANT: Pohlmann, Rainer
APPLICANT: Steiner, Sabine
APPLICANT: Mohr, Christine
APPLICANT: Wendland, Jürgen
APPLICANT: Knechtle, Philipp
APPLICANT: Repischung, Corinne
TITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSSPYI
TITLE OF INVENTION: AND USES THEREOF
NUMBER OF SEQUENCES: 1152
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 6239264artis Corporation
STREET: 3054 Cornwallis Road
CITY: Research Triangle Park
STATE: No. 6239264th Carolina
COUNTRY: USA
ZIP: 27709
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: CH 0016/97

```

1      FILING DATE: 31-DEC-1996
2
3      ATTORNEY/AGENT INFORMATION:
4
5      NAME: Meigs, J. Timothy
6
7      REGISTRATION NUMBER: 38,241
8
9      REFERENCE/DOCKET NUMBER: PE/5-30306/A/CGL1976
10
11     TELECOMMUNICATION INFORMATION:
12
13     TELEPHONE: 919-541-8587
14
15     TELEFAX: 919-541-8699
16
17     INFORMATION FOR SEQ ID NO: 288:
18
19     SEQUENCE CHARACTERISTICS:
20
21     LENGTH: 837 base pairs
22
23     TYPE: nucleic acid
24
25     STRANDEDNESS: single
26
27     TOPOLOGY: linear
28
29     MOLECULE TYPE: DNA (genomic)
30
31     ORIGINAL SOURCE:
32
33     ORGANISM: PAGI241RP
34
35     OS-08-998-416-288

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Query Match	17.2%;	Score 56.4;	DB 4;	Length 837;
Best Local Similarity	52.1%;	Pred. No. 0.07;		
Matches 149;	Conservative	0;	Mismatches 136;	Indels 1;
			Gaps	1

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QY	98	tgaatagatagttaataaataaagaatgtctcgggggaacttttgcgtttaaaag	157
Db	385	AATTTACAAFTTTTAAATTAATTTAACTTTATATATATAAAATATTTTAAATAAA	3369
QY	158	gaataataaataattagataaagctgtaaaataatctatttctaattgtt	217
Db	325	CAATTAAATTAATTAATTAATTAATTCGA-TAAATCTTTTAATTAATTAATTAAGAAAT	267
QY	218	aaaaattgatalaatgaaattgtaaaaaaaattcagggggagataaaatgaaaaa	2777
Db	266	AATTAATATCTAATTAATTTTAAATCACTAATTTAAATTTGAACATGACCTAATACTAT	207
QY	278	ttatttcaagtttaccgtaatttttgcgttttcaatgtttcttat	323
Db	206	TCAATTTAAATTAATTTTAAATTAATTAATTAATTAATTAATTAATTAATTAAT	161

RESULT 10
US-08-617-860B-33

GENERAL INFORMATION:
APPLICANT: Typfer, R., Bauror, J., Bothmann, H., Flisk, E.
APPLICANT: Hwicke-Grandpierre, C., Klein, B., Martini, N.,
APPLICANT: M iller, A., Schulte, W., Voeltz, M., Walek, J.,
APPLICANT: Schell, J.
TITLE OF INVENTION: Promoters
NUMBER OF SEQUENCES: 35
CORRESPONDENCE ADDRESS:
ADDRESSEE: Steinberg, Raskin & Davidson, P.C.
STREET: 1140 Avenue of the Americas

COMPUTER READABLE FORM:
MEDIUM TYPE: diskette, 3.50 inch, 1.4 Mb storage
COMPUTER: IBM compatible
OPERATING SYSTEM: DOS
SOFTWARE: Patentin Release #1.0, Version #1.25 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/617,860B
FILING DATE: 01-MAR-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/EP94/02950

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1      FILLING DATE: 05-SEP-1994
2      APPLICATION NUMBER: DE P4329951.2
3      FILING DATE: 04-SEP-1993
4      INFORMATION FOR SEQ ID NO: 33:
5          SEQUENCE CHARACTERISTICS:
6              LENGTH: 2750 Base pairs
7              TYPE: Nucleic acid
8              STRANDEDNESS: Double
9              TOPOLOGY: linear
10         MOLECULE TYPE: DNA (genomic)
11         HYPOTHEetical: NO
12         ANTI-SENSE: NO
13         ORIGINAL SOURCE:
14             ORGANISM: Cuphea lanceolata
15             IMMEDIATE SOURCE:
16             LIBRARY: genomic lambda FIX II
17             CLONE: Cl194
18     FEATURE:
19         NAME/KEY: Startcodon
20         LOCATION: 2637..2639
21     FEATURE:
22         NAME/KEY: CDS
23         LOCATION: 2637..2750
24     OS-08-617-860B-33

```

	Query Match	17.1%	Score 55.8	DB 3	Length 2750;
	Best Local Similarity	53.2%	Pred. No. 0.077;		
	Matches 143;	Conservative 0;	Mismatches 122;	Indels 4;	Gaps
OY	34	gtttaaataagaataagaataaaaaagatalatatcatatagctgaattataat	93		
Db	1505	GATTAAATTAATAAATAAGACTTAGTCGAATTTAATATATGACCAATTTTAACT	1566		
OY	94	tatagtaatagtagtiaataataaaaagtcttcgggagacacttttgttta	153		
Db	1565	ATTTTATTAGTAAAAAAAACGTAAAAAGATTTGTATTTGGTTTTCAATTTTCTCT	1624		
OY	154	aaaggaaatacataataaaattagataaaagtgtataaattc-----atttcctta	209		
Db	1625	CAAATGAAATTTATACCAAAATTAATATATTTAAATAAAAATTTAAATTTAAAGCATAAATTTGT	1684		
OY	210	aatttgcttaaaaattcgatataatgcgaatgtgtaaaaaaaatttcgggggagataaat	269		
Db	1685	AATTTTTTAATCTCGACTAATTTATTTTCGTTAATATTTAAAAATTTAGTTGAAA	1744		
OY	270	gaaaaaaataatttcctaagtttcgttaa	298		
Db	1745	AGGCATTAATATTCATTAACAATTTTGAA	1773		

RESULT 11
US-08-883-795A-36
; Sequence 36, Application US/08883795A

GENERAL INFORMATION:
APPLICANT: Delcuve, Genevieve
APPLICANT: Awang, Gregor
TITLE OF INVENTION: Recombinant DNA Molecules and Expression
TITLE OF INVENTION: Vectors for Tissue Plasminogen Activator
NUMBER OF SEQUENCES: 39
CORRESPONDENCE ADDRESS:
ADDRESSEE: BERSKIN & PARR
STREET: 40 King Street West

```

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25

```


CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/883,795A
FILING DATE: 27-JUN-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Graveille, Michelle
REGISTRATION NUMBER: 40,261
REFERENCE/DOCKET NUMBER: 7841-062
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 364-7311
TELEFAX: (416) 361-1398
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 665 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
IMMEDIATE SOURCE:
CLONE: Rh 32
US-08-883-795A-36

Query Match 17.0%; Score 55.6; DB 2; Length 665;
Best Local Similarity 50.38; Pred. No. 0.096;
Matches 163; Conservative 0; Mismatches 159; Indels 4; Gaps 1;

QY 1 attggagatcacttaaatcttagcagaagaatgtttaacgaataagaataaataa 60
DB 34 ATATTTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 93
QY 61 gatataatataatagctgaaaaattataatataatgataagatgtaataataa 120
DB 94 AATATTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 153
QY 121 aaagtgtcttcgggggacacttttcttcaaaaaggaataataaattagat 180
DB 154 AATATTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 213
QY 181 aaagtgaataataatttttttttttttttttttttttttttttttttttttt 236
DB 214 AATATTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 273
QY 237 atcgtaaaaaaattcaggggggaataataaagaaaaattttcaagttactgt 296
DB 274 AATGTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 333
QY 297 aattttatgtttcagtttttttttttttttttttttttttttttttttttt 324
DB 334 AATGTTTATTAATAAATTTATTAATAAATTTATTAATAAATTTATTAATAA 361

RESULT 12
US-08-446-855A-1/c
Sequence 1, Application US/08446855A
Patent No. 5849573
GENERAL INFORMATION:
APPLICANT: Stewart, Thomas S
APPLICANT: Flores, Maria V
APPLICANT: O'Sullivan, William J
TITLE OF INVENTION: Nucleotide sequence encoding carbamoyl
NUMBER OF SEQUENCES: 2
CORRESPONDENCE ADDRESS:
ADDRESSEE: Nixon & Vandierne PC
STREET: 1100 No. 5849573th Glebe Road, 8th Floor
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22201-4714
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.24
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/446,855A
FILING DATE: 06-Jul-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mitchard, Leonard C
REGISTRATION NUMBER: 29,009
REFERENCE/DOCKET NUMBER: 47-80
TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-816-4000
TELEFAX: 703-816-4100
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 8920 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic
US-08-446-855A-1

Query Match 16.88%; Score 54.8; DB 2; Length 8920;
Best Local Similarity 53.38; Pred. No. 0.099;
Matches 163; Conservative 0; Mismatches 137; Indels 6; Gaps 2;

QY 28 aagaatgtttaaatgaataagaataaagaalatalattatataatagctgaanaat 87
DB 8783 AAAAGAAATTTATACATTTTATGTTATATATTTTATTTTATTAATAAATTT 8724
QY 88 tataatataatgataagatagtttaataaataaagaatgtctcgggggacacttttg 147
DB 8723 ATTAATAATTAATAAATTTGTAATGTAATAAATGCAAAATTTGTTTACATATGACTG 8664
QY 148 tttaaaaaa---ggaataataaatttagataaagaatgtaaaataattttt 203
DB 8663 AATAATAAATTTGTAATATTAACAAATAATTTATTAATAAATAATCAATTAATAATTA 8604
QY 204 atttaatttgtaaaa--attgataataatgtaaaataaatttcaagggggga 261
DB 8603 TGACATATTTATTAATAAATTTTATTAATTTTAAGCTAATTAATTAATAATAA 8544
QY 262 atataaatgaaaaaattattcaagtttactgtaattttatgttttcaatgtttc 321
DB 8543 AT 8484
QY 322 attgtt 327
DB 8483 ACTTTT 8478

RESULT 13
US-09-150-741-1/c
Sequence 1, Application US/09150741
Patent No. 6183996
GENERAL INFORMATION:
APPLICANT: Stewart et al.
TITLE OF INVENTION: Nucleotide Sequence Encoding Carbamoyl Phosphate
PATENT NO. 6183996
FILE REFERENCE:
CURRENT APPLICATION NUMBER: US/09/150,741
CURRENT FILING DATE: 1998-09-10
EARLIER APPLICATION NUMBER: PL6380
EARLIER FILING DATE: 1997-12-16
EARLIER APPLICATION NUMBER: AU93/00617
EARLIER FILING DATE: 1993-12-02
EARLIER APPLICATION NUMBER: 08/446,855
EARLIER FILING DATE: 1995-07-06
NUMBER OF SEQ ID NOS: 15

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:27:08 ; Search time 222.28 Seconds
(without alignments)
1261.226 Million cell updates/sec

Title: US-09-531-438-3
Perfect score: 327
Sequence: 1 attgggatatcttaattt.....tttcattgttctattgtt 327

Scoring table: OLIGO_NUC
Gapop 60.0 , Capext 60.0

Searched: 930621 seqs, 428662619 residues

Word size: 0

Total number of hits satisfying chosen parameters: 989696

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

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- 11: /SIDS2/gcgdata/geneseq/geneseq/NA1990.DAT:*
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- 13: /SIDS2/gcgdata/geneseq/geneseq/NA1992.DAT:*
- 14: /SIDS2/gcgdata/geneseq/geneseq/NA1993.DAT:*
- 15: /SIDS2/gcgdata/geneseq/geneseq/NA1994.DAT:*
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- 17: /SIDS2/gcgdata/geneseq/geneseq/NA1996.DAT:*
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- 19: /SIDS2/gcgdata/geneseq/geneseq/NA1998.DAT:*
- 20: /SIDS2/gcgdata/geneseq/geneseq/NA1999.DAT:*
- 21: /SIDS2/gcgdata/geneseq/geneseq/NA2000.DAT:*
- 22: /SIDS2/gcgdata/geneseq/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	16	4.9	29	17	AAT42655
2	16	4.9	29	21	AAAS1200
3	16	4.9	32	20	AAZ27689
4	16	4.9	36	18	AAT97603
5	16	4.9	39	20	AAV90775
6	16	4.9	47	21	AAZ68379
7	15	4.6	20	21	AAA66183
8	15	4.6	24	22	AAH46804
9	15	4.6	27	17	AAT39439
10	15	4.6	27	19	AAV37457
11	15	4.6	30	14	AAQ43975

12	15	4.6	48	16	AAT25576	Human gene signatu
13	14	4.3	18	13	AAO20161	Cross-linking olig
14	14	4.3	18	13	AAO20160	Oligomer HSV724 fo
15	14	4.3	18	13	AAO30311	Oligomer HSV723 fo
16	14	4.3	18	13	AAO30310	Human C-IAP-1 mRNA
17	14	4.3	18	13	AAZ22162	Human ICGAP2 CPG 1
18	14	4.3	22	22	AAO1590	Human ICGAP2-5'-tr
19	14	4.3	22	22	AAO15643	Helicobacter pylori
20	14	4.3	26	19	AAV07952	Helicobacter pylori
21	14	4.3	26	19	AAV07922	Helicobacter pylori
22	14	4.3	27	19	AAV07937	Primer for amplify
23	14	4.3	29	17	AAT42655	N-terminal primer
24	14	4.3	29	21	AAAS1200	Human gene signatu
25	14	4.3	31	16	AAT52703	Nucleotide fragmen
26	14	4.3	31	19	AAV67854	PCR primer for Ver
27	14	4.3	32	20	AAZ27689	Shigella dysenteriae
28	14	4.3	36	18	AAT97603	Tomato spotted wil
29	14	4.3	36	22	AAO90606	Glycophorin antibo
30	14	4.3	37	15	AAO62952	Oligonucleotide us
31	14	4.3	45	22	AAE55449	Oligonucleotide TA
32	14	4.3	45	22	AAE55450	Oligonucleotide TA
33	14	4.3	45	22	AAO88874	Oligonucleotide TA
34	14	4.3	45	22	AAO88875	Human map-related
35	14	4.3	47	21	AAZ66366	Human map-related
36	14	4.3	47	21	AAZ67473	Human map-related
37	14	4.3	47	21	AAZ67533	Human map-related
38	14	4.3	47	21	AAZ67549	Human map-related
39	14	4.3	47	21	AAZ67813	Human map-related
40	14	4.3	50	21	AAAS8312	Human map-related
41	13	4.0	15	22	AAE48097	IGFBP3 oligonucleo
42	13	4.0	15	22	AAE48098	IGFBP3 oligonucleo
43	13	4.0	15	22	AAE48099	IGFBP3 oligonucleo
44	13	4.0	16	21	AAAS7758	Nucleotide sequenc
45	13	4.0	17	16	AAO92084	Renilla reniformis

ALIGNMENTS

RESULT 1

AAT42655 standard; DNA; 29 BP.

ID AAT42655:

AC AAT42655:

DT 25-FEB-1997 (first entry)

XX

DE Primer for amplifying verotoxin (VT-1) subunit A coding sequence.

XX

KW Verotoxin; Escherichia coli; enteric infection; diarrhoea; vaccine;

KM haemolytic uraemic syndrome; detection; ss.

XX

OS Synthetic

XX

PN WO9630043-A1

XX

PD 03-OCT-1996.

XX

PE 25-MAR-1996; 96WO-US04093.

XX

PR 24-MAR-1995; 95US-0410058.

XX

XX (OPHT-) OPHIDIAN PHARM INC.

XX

XX Carroll SB, Padhye NV, Stafford DC;

XX

XX WPI; 1996-505779/50.

XX

XX Compos. contrg. neutralising antitoxin against E.coli vero-toxin -

XX PT used to treat intoxicated individuals, and as a prophylactic against

XX PT diarrhoeal disease or extra-intestinal complications of E.coli

XX PT infection

PS Example 6; Page 58; 101pp; English.

CC Compositions containing neutralising antitoxin against one or more E.

CC coli verotoxin (VT) can be used to treat intoxicated adults and

CC children with enteric bacterial infections. They may also be used as

CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the

CC development of extra-intestinal complications of E.coli infection,

CC especially haemolytic uraemic syndrome. The antitoxin can also be

CC used to detect E. coli VT in a sample. The VT is recombinant,

CC preferably a fusion protein containing a non-VT protein sequence and

CC part of the E.coli VT1 or VT2 sequence. Two primers (AA12655,

CC AA12656) were used to amplify the verotoxin VT-1 A subunit coding

CC sequence and add a histidine tag coding sequence to the subunit

CC sequence. Two primers (AA12655, AA12658) were used to amplify the

CC verotoxin VT-1 A and B subunits and add a histidine tag coding

CC sequence to the subunit sequences.

CC SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

XX

Query Match 4.9%; Score 16; DB 17; Length 29;

Best Local Similarity 100.0%; Pred. No. 1e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204

Db 9 aaataattatttta 24

RESULT 2

AA12600

AA12600 standard; DNA; 29 BP.

XX

AC AA12600;

XX

DT 26-SEP-2000 (first entry)

XX

DE N-terminal primer for E. coli verotoxin 1 subunit A gene.

XX

KM VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;

KM recombinant production; screening; dairy; anti-bacterial; vaccine;

KM primer; polyhistidine; ss.

XX

OS Escherichia coli.

OS Synthetic.

XX

PN US6080400-A.

XX

PD 27-JUN-2000.

XX

PF 13-MAR-1997; 97US-0816977.

XX

PR 24-MAR-1995; 95US-0410058.

XX

PA (OPHI-) OPHIDIAN PHARM INC.

XX

PI Williams JA, Byrne LM;

XX

DR WPI; 2000-451195/39.

XX

PT Bacterial cell for recombinantly expressing bacterial toxins in large

PT quantities useful for immunization and treatment of bacterial

PT infections, comprises expression vector encoding bacterial toxin

XX

PS Example 6; Column 83; 83pp; English.

XX

CC E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into

CC pET-23b, designed to allow expression of the native proteins containing

CC C-terminal polyhistidine tags. The VT-1 and VT-2 genes were engineered

CC to convert the signal sequence methionine codon into a NdeI site to

CC allow cloning of the amplified genes into the vector without addition of

CC vector-encoded amino acids. The C-terminal primers comprises the

CC C-terminal 7 codons of each gene fused to the sequence CTCGAGCC, in order

CC to add the polyhistidine tag. The primers delete the native stop codons,

CC and when cloned into pET-23 add a C-terminal extension of Leu-Glu-(His)₆.

CC VT B chains are small proteins (approximately 8 kDa), so use of a small

CC affinity tag was preferred (i.e. polyhistidine). A polyhistidine affinity

CC tag facilitates single step affinity purification of subunits from

CC periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A

CC and VT-2 A chains, expression of maltose binding protein (MBP) fused

CC subunits was undertaken. Due to the toxicity of the VT-2 B subunit,

CC strict uninduced promoter control is necessary to permit cell viability.

CC Bacterial host cells expressing a recombinant expression vector encoding

CC a polyhistidine affinity tag and a portion of the VT-2 B chain are

CC claimed. The vector is chosen from PET24hisVT2BL⁺, PET24hisVT2BL⁻ and

CC PET24VT2B, where "L⁺" indicates that the vector encodes the preprotein

CC form of the protein and "L⁻" indicates that the vector encodes the mature

CC form of the protein. The bacterial cell is capable of expressing large

CC quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing

CC non-mammals and for detecting bacterial toxins in environmental samples

CC including soil, water, industrial samples, biological samples and samples

CC obtained from food and dairy processing instruments.

CC SQ Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

XX

Query Match 4.9%; Score 16; DB 21; Length 29;

Best Local Similarity 100.0%; Pred. No. 1e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204

Db 9 aaataattatttta 24

RESULT 3

AA227689

AA227689 standard; DNA; 32 BP.

XX

AC AA227689;

XX

DT 22-DEC-1999 (first entry)

XX

DE PCR primer for Verotoxin gene.

XX

KM Verotoxin; VT1; VT2; detection; PCR primer; ss.

XX

OS Synthetic.

OS Escherichia coli.

XX

PN JP11243996-A.

XX

PD 14-SEP-1999.

XX

PF 27-FEB-1998; 98JP-0047677.

XX

PR 27-FEB-1998; 98JP-0047677.

XX

PA (TOYM) TOYOCO KK.

XX

DR WPI; 1999-603716/52.

XX

PT An oligonucleotide for amplification of verotoxin - useful in the

PT detection of inactivated verotoxin gene by transfer of a foreign DNA

PT fragment

XX

PS Claim 11; Page 9; 10pp; Japanese.

XX

CC This sequence represents a PCR primer of the invention. The primer is

CC used for amplification of the E. coli verotoxin (VT) gene. The

CC oligonucleotide is useful for detection of inactivated VT gene by

CC transfer of a foreign DNA fragment. Simple, rapid and specific

CC amplification of VT gene from environmental factors is achieved using the

CC oligonucleotide of the invention.

XX

SQ Sequence 32 BP; 12 A; 2 C; 4 G; 14 T; 0 other;

Query Match 4.9%; Score 16; DB 20; Length 32;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||
 Db 4 aaataattatttta 19

RESULT 4

AAV97603
 ID AAT97603 standard; DNA; 36 BP.
 XX AAT97603;
 XX

30-APR-1998 (first entry)

XX Shigella dysenteriae delta-stx_A allele PCR primer 13.

XX Delta-virg allele; delta-guab-A allele; PCR; amplification; primer;
 KW delta-stx_A allele; shigellosis; vaccine; ss.

XX Synthetic.
 OS Shigella dysenteriae.

XX W09737685-A1.

XX 16-OCT-1997.

XX 09-APR-1997; 97WO-US05954.

XX 09-APR-1996; 96US-0629600.

XX (UYMA-) UNIV MARYLAND BALTIMORE.

XX Levine MM, Noriega FR;

XX WPI: 1997-512417/47.

XX Shigella mutants with mutation in guab-A - used in vaccines against
 Shigellosis

XX Example 6; Page 57; 94pp; English.

XX This is a PCR primer used in the amplification of the Shigella
 CC dysenteriae 1 delta-stx_A allele. The delta-stx_A allele was integrated
 CC into delta-guab-A of delta-guab-A, delta-virg S. dysenteriae 1, which
 CC inactivated the shiga toxin of this strain. The mutant can be used in
 CC the preparation of vaccines such as, a live vector vaccine comprising
 CC a Shigella mutant, (which encodes and expresses a foreign vaccine
 CC comprising the Shigella mutant (which also contains a plasmid which
 CC encodes and expresses a foreign antigen in a eukaryotic cell). The
 CC vaccines can be used against Shigellosis.

XX Sequence 36 BP; 11 A; 3 C; 10 G; 12 T; 0 other;

Query Match 4.9%; Score 16; DB 18; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204
 |||||
 Db 14 aaataattatttta 29

RESULT 5

AAV90775
 ID AAV90775 standard; DNA; 39 BP.
 XX

AC AAV90775;
 XX 18-FEB-1999 (first entry)
 DT
 XX Primer Y104F.

XX Antigen; immunogenic cluster family; vaccine; gastritis; diagnosis;
 KW peptic ulcer; gastric adenocarcinoma; gastric lymphoma; primer; ss.
 XX

OS Synthetic.

XX W09849314-A2.

XX 05-NOV-1998.

XX 27-APR-1998; 98WO-US08487.

XX 14-OCT-1997; 97US-0061958.

XX 25-APR-1997; 97US-0045107.

XX (GENE-) GENELABS TECHNOLOGIES INC.

XX Chow TP, Fry KE, Lim MY, McAttee CP;

XX WPI: 1999-009433/01.

XX New Helicobacter pylori antigens and related nucleic acid sequences
 PT - useful in serological diagnosis and protective vaccines, providing
 PT long-lasting immune response

XX Claim Disclosure; Page 194; 402pp; English.

XX The specification, which describes Helicobacter pylori antigenic
 CC proteins that are characterised by immunoreactivity with
 CC H. pylori-positive antisera. The specification also describes 69
 CC previously unrecognised immunogenic cluster families. H. pylori
 CC antigens are used to detect H. pylori-specific antibodies, for
 CC diagnosing infection or to confirm eradication of infection, and
 CC in vaccines to protect against H. pylori infection and related
 CC diseases (gastritis, peptic ulcer, gastric adenocarcinoma/lymphoma).
 CC The present primer is used in the course of the invention.

XX Sequence 39 BP; 15 A; 8 C; 7 G; 9 T; 0 other;

Query Match 4.9%; Score 16; DB 20; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 atacttaatttagc 23
 |||||
 Db 23 atacttaatttagc 38

RESULT 6

AAZ68379
 ID AAZ68379 standard; DNA; 47 BP.

XX AAZ68379;

XX 10-SEP-2001 (first entry)

XX Human map-related biallelic marker SEQ ID NO:2726.

XX Human genome; biallelic marker; high density disequilibrium map;
 KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
 KW haplotyping; hybridisation; identification; characterisation;
 KW diagnosis; single nucleotide polymorphism; SNP; ds.

XX Homo sapiens.

XX Key Location/Qualifiers
 FH variation replace(24,A)
 FT

```
FT      /*tag= a
XX      /standard_name= "single nucleotide polymorphism"
XX      W09954500-A2.
XX      PD
XX      28-OCT-1999.
XX      PF
XX      21-APR-1999; 99WO-IB00822.
XX      PR
XX      21-APR-1998; 98US-0082614.
XX      PR
XX      23-NOV-1998; 98US-0109732.
XX      PA
XX      (GEST ) GENSET.
XX      PI
XX      Cohen D, Blumenfeld M, Chumakov I;
XX      WPI: 2000-013267/01.
XX      DR
XX      WPI: 2000-013267/01.
XX      PT
XX      Novel biallelic markers used to construct a high density disequilibrium
XX      map of the human genome -
XX      PS
XX      Claim 3; Page 812; 2745pp; English.
XX      CC
XX      AA265654 to AA269578 represent human biallelic markers from the present
XX      CC
XX      invention, which contain a polymorphic base at position 24 of their
XX      CC
XX      nucleotide sequences. AA269579 to AA277440 represent amplification
XX      CC
XX      primers for the biallelic markers. The biallelic markers of the
XX      CC
XX      invention have a variety of uses: they can be used for high density
XX      CC
XX      mapping of the human genome, and in complex association studies and
XX      CC
XX      haplotyping studies which are useful in determining the genetic basis
XX      CC
XX      for disease states. Compositions and methods of the invention can also
XX      CC
XX      be useful for the identification of the targets for the development of
XX      CC
XX      pharmaceutical agents and diagnostic methods, as well as the
XX      CC
XX      characterisation of the differential efficacious responses to and side
XX      CC
XX      effects from pharmaceutical agents acting on a disease as well as other
XX      CC
XX      treatment.
XX      CC
XX      N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
XX      CC
XX      and 3367, are not actually given a sequence in the Sequence Listing
XX      CC
XX      from the present invention.
XX      SQ
XX      Sequence 47 BP; 24 A; 5 C; 6 G; 12 T; 0 other;

Query Match      4.9%; Score 16; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 9.8e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      147 gtttaaaaggaaga 162
        |||||||||||||||
DB      13 gtttaaaaggaaga 28

RESULT 7
ID      AAA66183 standard; DNA; 20 BP.
XX      AC
XX      AAA66183;
XX      DT
XX      09-OCT-2000 (first entry)
XX      DE
XX      Dog genomic marker oligonucleotide sequence SEQ ID NO:45.
XX      KW
XX      Dog; genome; genomic marker; radiation hybrid map; identification;
XX      KW
XX      chromosome location; gene marker; polymorphic microsatellite marker;
XX      KW
XX      phenotype; behaviour; pedigree; ss.
XX      OS
XX      Canis familiaris.
XX      PI
XX      WO200029615-A2.
XX      PD
XX      25-MAY-2000.
XX      PF
XX      15-NOV-1999; 99WO-IB01907.
```

```
XX      13-NOV-1998; 98US-0108193.
XX      PR
XX      (CNRS ) CNRS CENT NAT RECH SCI.
XX      PA
XX      Gallbert F, Andre C;
XX      PI
XX      WPI: 2000-387821/33.
XX      DR
XX      WPI: 2000-387821/33.
XX      PT
XX      New radiation hybrid map of the dog, Canine familiaris, genome, useful
XX      PT
XX      for e.g. identifying genes implicated in phenotypic and behavioral
XX      PT
XX      traits or in genetic diseases and for studying dog pedigrees -
XX      PS
XX      Claim 1; Page 55; 87pp; English.
XX      CC
XX      The present invention describes a radiation hybrid map of the dog
XX      CC
XX      (Canine familiaris) genome comprising the genome location of a marker
XX      CC
XX      selected from AA66139 to AA66942. The radiation hybrid map is useful
XX      CC
XX      for identifying and localising dog genes, since it covers approximately
XX      CC
XX      80 % of the dog genome and provides a dense map integrating different
XX      CC
XX      types (i.e. Type I and Type II) of markers. The map and the dog genome
XX      CC
XX      markers (or complementary sequences) are especially useful to identify
XX      CC
XX      genes responsible for phenotypic and behavioural traits in dogs, to
XX      CC
XX      identify morbid genes, to analyse diseases and identify implicated genes
XX      CC
XX      in such diseases and their alleles, and to study dog pedigrees. They
XX      CC
XX      may also be useful for isolating corresponding human gene sequences
XX      CC
XX      e.g. genes involved in genetic diseases.
XX      SQ
XX      Sequence 20 BP; 14 A; 1 C; 2 G; 3 T; 0 other;

Query Match      4.6%; Score 15; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      261 aatataatgaaaa 275
        |||||||||||||||
DB      1 aatataatgaaaa 15

RESULT 8
ID      AAH46804 standard; DNA; 24 BP.
XX      AC
XX      AAH46804;
XX      DT
XX      19-SEP-2001 (first entry)
XX      DE
XX      Human high motility group protein family 11 cDNA PCR primer #2.
XX      KW
XX      Human; high motility group protein family 11; cancer; haemopathy;
XX      KW
XX      HIV infection; immune disease; inflammation; gene therapy;
XX      KW
XX      PCR primer; ss.
XX      OS
XX      Homo sapiens.
XX      PI
XX      WO200147967-A1.
XX      PD
XX      05-JUL-2001.
XX      PF
XX      18-DEC-2000; 2000WO-CN00595.
XX      PR
XX      23-DEC-1999; 99CN-0125721.
XX      PA
XX      (UYFU-) UNIV FUDAN.
XX      PA
XX      (SHAN-) SHANGHAI BIO DOOR GENE TECHNOLOGY LTD.
XX      PI
XX      Mao Y, Xie Y;
XX      PD
XX      WPI: 2001-418226/44.
XX      DR
XX      WPI: 2001-418226/44.
XX      PT
XX      High motility group protein family 11 and encoded polynucleotide,
XX      PT
XX      applicable in diagnosis and treatment of cancer, haemopathy, HIV
```


PT	infection, immunological diseases and various inflammation
XX	
PS	Example 3; Page 17; 39pp; Chinese.
CC	The present invention provides the protein and coding sequences of the
CC	human high motility group protein family 11. The sequences are useful in
CC	the treatment of cancer, haemopathy, HIV infection, immune diseases and
CC	inflammation. The present sequence is a PCR primer for the coding
CC	sequence of the invention.
XX	
SO	Sequence 24 BP; 8 A; 3 C; 0 G; 13 T; 0 other;
OY	
Db	Query Match 4.6%; Score 15; DB 22; Length 24; Best Local Similarity 100.0%; Pred. No. 2.6e+03; Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps
	193 taattatttttattt 207 2 taactatttttattt 16
RESULT	9
ID	AAT39439 standard; RNA; 27 BP.
XX	
AC	AAT39439;
DT	13-NOV-1996 (first entry)
XX	
DE	Hel-N2 selected sequence, e-13.
DE	
KW	Human, neuron-specific protein; Hel-N1, 3'-UTR instability sequence;
KM	paraneoplastic sensory neuropathy; oncoprotein; lymphokine; rat; elav;
KM	RNA recognition motif; RRM; Drosophila; cellular growth; localisation;
KW	instability; transistability; neurons; autoimmune protein; PE; PCD; PSN;
KM	central nervous system; cancer; paraneoplastic cerebellar degeneration;
KW	paraneoplastic encephalomyelitis; RNP-1 octamer sequence;
KW	humoral elav-like neuronal protein-1; ss.
XX	
OS	Synthetic.
FH	
FT	Key Location/Qualifiers
FT	misc_RNA 15..23
FT	/*tag= a
FT	/note= "Consensus sequence"
XX	
PN	US5525495-A.
XX	
PD	11-JUN-1996.
XX	
PE	11-MAY-1992; 92US-0881075.
XX	
PR	15-SEP-1993; 93US-0120827.
XX	
PR	11-MAY-1992; 92US-0881075.
XX	
PA	(UYDU-) UNIV DUKE.
PI	
PI	Gao F, Keene JD, Levine T;
DR	WPJ: 1996-286398/29.
XX	
PT	Prod'n of cDNA library for related proteins - by screening total cell
XX	mRNA with RNA-binding protein Hel-N1 or Hel-N2
PS	Disclosure; Fig 11; 66pp; English.
CC	The sequences given in AAT39427-46 represent random oligonucleotides
CC	which were isolated due to their ability to bind to the human neuron-
CC	-specific protein, Hel-N2. These sequences contain short stretches
CC	of uridylylate residues interspersed with other residues. These U-
CC	-rich regions share homology with the 3'-UTR instability sequences
CC	that are found in mRNA's. Instability sequences are target elements

CC		which reside in the 3'-non-coding regions of mRNA's which encode
CC		oncoproteins and lymphokines. Hel-N2 is a deleted form of Hel-N1
CC		in which the residues 239-251 have been deleted. This protein is
CC		expressed in medulloblastoma tumor cells and is not found in whole
CC		human brain. A small amount of Hel-N2 is also found in fetal brain,
CC		which may indicate a correlation with rapid growth. Hel-N1 cDNA was
CC		isolated by probing for rat and human elay counterparts using degenerate
CC		primers designed to simulate the RNP-1 octamer sequence present in two
CC		of the three RNA recognition motifs (RRW's) of Drosophila elav. Hel-N1
CC		was found to contain 3 RRW's, where the third one (see also AA000244) is
CC		sufficient for mRNA 3'-UTR binding activity. Full length Hel-N1, when
CC		transfected into a cell, causes cellular growth to cease, however, if
CC		just the third binding domain is transfected into cells, the cells
CC		undergo rapid growth. Hel-N1 binds as a multimer along the mRNA,
CC		presumably enhancing its localisation, instability and/or regulating it
CC		translatability and/or deadenylation it. This protein may be
CC		responsible for the growth cessation of neurons. Hel-N1 is an autolimmune
CC		protein in certain patients who show central nervous system
CC		manifestations of cancer called paraneoplastic cerebellar degeneration
CC		(PCD), paraneoplastic encephalomyelitis (PE) or paraneoplastic sensory
CC		neuropathy (PSN).
CC		
SQ		Sequence 27 BP; 7 A; 2 C; 2 G; 16 U; 0 other;
OY	197 tatatttatttaaa 211	
Dd	12 uauuuuuuuuuuaa 26	
RESULT 10		
AAV37457		
ID	AAV37457 standard; RNA; 27 BP.	
XX		
AC	AAV37457;	
XX		
DT	07-SEP-1998 (first entry)	
DE		
XX		
KM	Human Hel-N2 selected RNA sequence e-13.	
XX		
KW	Growth regulatory protein; Hel-N2; oncogene; cytokine; lymphokine;	
XX	chromosome mapping; human; functionally related protein; ss.	
OS	Homo sapiens.	
PN	US5773246-A.	
PD	30-JUN-1998.	
Pf	07-JUN-1995; 95US-0478675.	
PR	15-SEP-1993; 93US-0120827.	
PR	11-MAY-1992; 92US-0881075.	
PR	07-JUN-1995; 95US-0478675.	
PA	(GAOF/) GAO F.	
PA	(KEEN/) KEENE J D.	
PA	(LEVI/) LEVINE T.	
PI	Gao F, Keene JD, Levine T;	
DR	WPI; 1998-387003/33.	
PT	Use of proteins which bind RNA - for obtaining a cDNA library	
PT	containing members encoding structurally or functionally related	
PT	proteins from total cell mRNA.	
PS	Disclosure; Fig 11; 67pp; English.	
XX		

Query Match 4.6%; Score 15; DB 16; Length 48;
 Best Local Similarity 100.0%; Pred. No. 2.5e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 159 aaataataataaa 173
 |||||
 Db 17 aaataataataaa 31

RESULT 13

AAQ20161
 ID AAQ20161 standard; DNA; 18 BP.

AC AAQ20161;

XX 01-APR-1992 (first entry)

DE Cross-linking oligomer 724 to target Herpes Simplex Virus I.

XX deoxyribonucleic acid; major groove; HSV;

KW Inverted polarity region; covalent cross-linking group; ss.

XX Synthetic.

Key location/Qualifiers

FT modified_base

FT /*tag= a

FT /mod_base= OTHER

FT modified_base

FT /*tag= b

FT /mod_base= OTHER

FT modified_base

FT /*tag= c

FT /mod_base= OTHER

FT modified_base

FT /*tag= d

FT /mod_base= OTHER

FT modified_base

FT /*tag= e

FT /mod_base= OTHER

FT modified_base

FT /*tag= f

FT /mod_base= OTHER

FT modified_base

FT /*tag= g

FT /mod_base= OTHER

FT modified_base

FT /*tag= h

FT /mod_base= OTHER

FT modified_base

FT /*tag= i

FT /mod_base= OTHER

FT modified_base

FT /*tag= j

FT /mod_base= OTHER

FT modified_base

FT /*tag= k

FT /mod_base= OTHER

FT modified_base

FT /*tag= l

FT /mod_base= OTHER

FT modified_base

FT /*tag= m

FT /mod_base= OTHER

FT modified_base

FT /*tag= n

FT /mod_base= OTHER

FT modified_base

FT /*tag= o

FT /mod_base= OTHER

FT modified_base

FT /*tag= p

FT /mod_base= OTHER

FT modified_base

FT /note= "N-methyl-8-oxo-2'-deoxyadenine"

FT /*tag= m

FT /mod_base= OTHER

FT /note= "N-methyl-8-oxo-2'-deoxyadenine"

FT modified_base

FT /*tag= n

FT /mod_base= OTHER

FT /note= "N-methyl-8-oxo-2'-deoxyadenine"

FT modified_base

FT /*tag= o

FT /mod_base= OTHER

FT modified_base

FT /*tag= p

FT /mod_base= OTHER

FT modified_base

FT /*tag= q

FT /mod_base= OTHER

FT modified_base

FT /*tag= r

FT /mod_base= OTHER

FT modified_base

FT /*tag= s

FT /mod_base= OTHER

FT modified_base

FT /*tag= t

FT /mod_base= OTHER

FT modified_base

FT /*tag= u

FT /mod_base= OTHER

FT modified_base

FT /*tag= v

FT /mod_base= OTHER

FT modified_base

FT /*tag= w

FT /mod_base= OTHER

FT modified_base

FT /*tag= x

FT /mod_base= OTHER

FT modified_base

FT /*tag= y

FT /mod_base= OTHER

FT modified_base

FT /*tag= z

FT /mod_base= OTHER

FT modified_base

FT /*tag= AA

FT /mod_base= OTHER

FT modified_base

FT /*tag= AB

FT /mod_base= OTHER

FT modified_base

FT /*tag= AC

FT /mod_base= OTHER

FT modified_base

FT /*tag= AD

FT /mod_base= OTHER

FT modified_base

FT /*tag= AE

FT /mod_base= OTHER

FT modified_base

FT /*tag= AF

FT /mod_base= OTHER

FT modified_base

FT /*tag= AG

FT /mod_base= OTHER

FT modified_base

FT /*tag= AH

FT /mod_base= OTHER

FT modified_base

FT /*tag= AI

FT /mod_base= OTHER

FT modified_base

Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;

Query Match 4.3%; Score 14; DB 13; Length 18;
 Best Local Similarity 100.0%; Pred. No. 6.9e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 159 aaataataataaa 172
 |||||
 Db 2 aaataataataaa 15

RESULT 14

AAQ20160
 ID AAQ20160 standard; DNA; 18 BP.

AC AAQ20160;

XX 01-APR-1992 (first entry)

DE Cross-linking oligomer 723 to target Herpes Simplex Virus I.

XX deoxyribonucleic acid; major groove; HSV;

KW Inverted polarity region; covalent cross-linking group; ss.

XX Synthetic.

Key location/Qualifiers

FT modified_base

FT /*tag= a

FT /mod_base= OTHER

FT modified_base

FT /*tag= b

FT /mod_base= OTHER

FT modified_base

FT /*tag= c

FT /mod_base= OTHER

FT modified_base

FT /*tag= d

FT modified_base /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 3 /tag= c
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 4 /tag= d
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 5 /tag= e
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 7 /tag= f
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 9 /tag= g
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 10 /tag= h
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 11 /tag= i
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 12.18 /tag= j
FT /label= inverted_polarity_region
FT /note= "see comments"
FT 13 /tag= k
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 14 /tag= l
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 15 /tag= m
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT 17 /tag= n
FT /mod_base= OTHER
FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
FT W09118997-A.
FT 12-DEC-1991.
FT 24-MAY-1991; 91WO-1003680.
FT 14-JAN-1991; 91US-0640654.
FT 25-MAY-1990; 90US-0529346.
FT (GILE-) GILEAD SCIE INC.
FT PA
FT XX
FT PI
FT Matreucci MD, Krawczyk S;
FT WPI; 1992-007480/01.
FT
FT New sequence-specific non-photo-activated crosslinking agents -
FT bind to the major groove of duplex DNA and are esp. useful for
FT treating latent infections e.g. HIV
FT
FT Example 4; Page 29; 42pp; English.
FT
FT This oligomer contains an inverted polarity region formed from an
FT o-xyloso dimer synthon. Residues 11 and 12 are linked via an

CC o-xyloso group (i.e. nucleotides that have xylose sugar linked via
CC the o-xyloso ring). The sequence is designed to target the Herpes
CC Simplex virus I beginning at nucleotide 10996 and to covalently
CC cross-link to it. See also AAQ20151-020161.
XX
SQ Sequence 18 BP; 13 A; 0 C; 0 G; 5 T; 0 other;

Query Match 4.3%; Score 14; DB 13; Length 18;
Best Local Similarity 100.0%; Pred. No. 6.9e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 159 aaatataataa 172
|||||
Db 2 aaatataataa 15

RESULT 15
AAQ30311
ID AAQ30311 standard; DNA; 18 BP.
XX
AC AAQ30311;
XX
XX 07-DEC-1992 (first entry)
DT
DE Oligomer HSV724 for forming triplex with HSV target duplex.
XX
KM Herpes simplex virus I; AIDS; modified; HIV; RSV; malignancy;
KM hepatitis; inflammation; ss.
OS
XX Synthetic.
FH
FH Key Location/Qualifiers
FT 1 /tag= a
FT /mod_base= OTHER
FT /note= "OTHER= N4 N4 ethanocytosine"
FT 2 /tag= b
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 3 /tag= c
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 4 /tag= d
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 5 /tag= e
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 7 /tag= f
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 9 /tag= g
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 10 /tag= h
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 11 /tag= i
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT 13 /tag= j
FT /mod_base= OTHER
FT /note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"
FT modified_base

FT		modified_base	14	/tag= k	OTHER
FT				/mod_base=	OTHER
FT		modified_base	15	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT				/tag= l	
FT		modified_base	17	/mod_base= OTHER	
FT				/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT				/tag= m	
FT				/mod_base= OTHER	
FT		misc_feature	12..18	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"	
FT				/tag= n	
FT				/label= Inverted.polarity_region	
FT		misc_feature	11..12	/note= "see comments"	
FT				/tag= o	
FT				/note= "O'-xylosio dimer synthon linkage"	
XX					
PM		W09209705-A.			
XX					
PD		11-JUN-1992.			
XX					
PE		25-NOV-1991;	91WO-US08811.		
XX					
PR		23-NOV-1990;	90US-0617907.		
PR		18-JAN-1991;	91US-0643382.		
PR		08-APR-1991;	91US-0683420.		
PR		17-APR-1991;	91US-0686544.		
PR		17-APR-1991;	91US-0686546.		
PR		17-APR-1991;	91US-0686547.		
PR		27-SEP-1991;	91US-0766733.		
XX					
PA		(GILE-) GILEAD SCI INC.			
XX					
PI		Froehner B, Krawczyk S, Matteucci MD, Malligan J;			
XX					
DR		WPI: 1992-217083/26.			
XX					
PT		New oligomers contg. modified bases - which form a triplex with			
PT		G-C doublet in a DNA duplex, for treating and diagnosing HIV,			
PT		hepatitis, herpes, malignancy and inflammation			
XX					
PS		Claim 12; Page 67; 77pp; English.			
XX					
CC		The synthetic oligomer is capable of forming a triplex at			
CC		physiological pH with a purine rich target sequence by coupling			
CC		into the major groove of the duplex. The specific target sequence			
CC		of this oligomer is a herpes simplex virus I duplex beginning at			
CC		nucleotide 10996 contg. a purine-rich region concentrated on			
CC		one chain of the duplex. The oligomer, and others like it are useful			
CC		in diagnosis and therapy of diseases characterised by specific DNA			
CC		duplex targets, e.g. respiratory syncytial virus, HIV, hepatitis,			
CC		herpes, malignant tumours and inflammation. The triple helices form			
CC		under mild conditions thus assays may be carried out without			
CC		subjecting the test specimen to harsh conditions. The oligomer			
CC		contains an inverted polarity region formed from an O'-xylosio			
CC		dimer synthon. The linking gp. is O'-xylosio (nucleotides have the 3'			
CC		positions of xylose sugars linked via the O'-xylene ring). Two			
CC		nucleotides are coupled through a xylene residue to form the dimer			
CC		synthon. This additional modifications may render the oligomer stable			
CC		to nuclease activity. The oligomer is able to inhibit gene expression,			
CC		as verified by in vitro systems.			
CC		See also AAQ25452-25501 and AAQ30226-448.			
XX					
SQ		Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;			
		Query Match	4.3%; Score 14; DB 13; Length 18;		
		Best Local Similarity	100.0%; Pred. NO. 6.9e+03;		
		Matches	14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		

QY		159	aaataataataaa	172	
Db		2	aaaataataataaa	15	
RESULT	16				
ID	AAQ30310				
XX	AAQ30310 standard; DNA; 18 BP.				
AC	AAQ30310;				
XX	07-DEC-1992 (first entry)				
DE	Oligomer HSV723 for forming triplex with HSV target duplex.				
XX					
KW	Herpes simplex virus I; AIDS; modified; HIV; RSV; HPV; malignan				
KW	hepatitis; inflammation; ss.				
XX					
OS	Synthetic..				
XX					
FH	Key				
FT	modified_base				
FT	Location/Qualifiers				
FT	1				
FT	/tag= a				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	2				
FT	/tag= b				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	3				
FT	/tag= c				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	4				
FT	/tag= d				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	5				
FT	/tag= e				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	7				
FT	/tag= f				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	9				
FT	/tag= g				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	10				
FT	/tag= h				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	11				
FT	/tag= i				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	13				
FT	/tag= j				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	14				
FT	/tag= k				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	15				
FT	/tag= l				
FT	/mod_base= OTHER				
FT	/note= "OTHER= N6 methyl-8-oxo 2' deoxyadenine"				
FT	17				
FT	/tag= m				

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FT misc-feature 12..18
FT /tag= n
FT /label= inverted_polarity_region
FT /note= "see comments"
FT misc-feature 11..12
FT /tag= 0
FT /note= "o-xylosa dimer synthon linkage"
PN WO9209705-A.
XX 11-JUN-1992.
XX
XX 25-NOV-1991; 91WO-US08811.
XX
XX 23-NOV-1990; 90US-0617907.
XX 18-JAN-1991; 91US-0643382.
XX 08-APR-1991; 91US-0683420.
XX 17-APR-1991; 91US-0686544.
XX 17-APR-1991; 91US-0686546.
XX 17-APR-1991; 91US-0686547.
XX 27-SEP-1991; 91US-0766733.
XX
XX (GILE-) GILEAD SCI INC.
XX
XX Froehner B, Krawczyk S, Matteucci MD, Milligan J;
XX WPI, 1992-217083/26.
XX
XX New oligomers contg. modified bases - which form a triplex with
XX G-C doublet in a DNA duplex, for treating and diagnosing HIV,
XX hepatitis, herpes, malignancy and inflammation
XX
XX Claim 12; Page 67; 77pp; English.
XX
XX The synthetic oligomer is capable of forming a triplex at
XX physiological pH with a purine rich target sequence by coupling
XX into the major groove of the duplex. The specific target sequence
XX of this oligomer is a herpes simplex virus I duplex beginning at
XX nucleotide 10996 contg. a purine-rich region concentrated on
XX one chain of the duplex. The oligomer, and others like it are useful
XX in diagnosis and therapy of diseases characterised by specific DNA
XX duplex targets, e.g. respiratory syncytial virus, HIV, hepatitis,
XX herpes, malignant tumours and inflammation. The triple helices form
XX under mild conditions thus assays may be carried out without
XX subjecting the test specimen to harsh conditions. The oligomer
XX contains an inverted polarity region formed from an o-xylosa
XX dimer synthon. The linking gp. is o-xylosa (nucleotides have the 3'
XX positions of xylose sugars linked via the o-xylyene ring). Two
XX nucleotides are coupled through a xylyene residue to form the dimer
XX synthon. This additional modifications may render the oligomer stable
XX to nuclease activity. The oligomer is able to inhibit gene expression,
XX as verified by in vitro systems.
XX See also AAQ25452-25501 and AAQ30226-448.
XX
XX Sequence 18 BP; 13 A; 0 C; 0 G; 5 T; 0 other;
XX
XX
XX Query Match 4.3%; Score 14; DB 13; Length 18;
XX Best Local Similarity 100.0%; Pred. No. 6.9e+03;
XX Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 159 aaataataataaa 172
XX |||||
XX Db 2 aaataataataaa 15
XX
XX RESULT 17
XX AA22162/C
XX ID AA22162 standard; DNA; 18 BP.
XX
XX AC AA22162;
XX
XX 26-NOV-1999 (first entry)
```

```
XX
XX Human c-IAP-1 mRNA inhibiting antisense oligo ISIS #23344.
XX DE
XX Cellular inhibitor of Apoptosis-1; antisense; diagnostic; therapeutic;
XX KM c-IAP-1; prophylaxis; infection; inflammation; tumor formation; ss.
XX KM
XX Synthetic.
XX OS Homo sapiens.
XX OS
XX US5958772-A.
XX
XX 28-SEP-1999.
XX
XX 03-DEC-1998; 98US-0205204.
XX
XX 03-DEC-1998; 98US-0205204.
XX
XX (ISIS-) ISIS PHARM INC.
XX
XX Bennett CF, Cowse LM, Ackermann EJ;
XX WPI, 1999-561047/47.
XX
XX Antisense compounds complementary to Cellular Inhibitor of Apoptosis-1
XX useful for e.g. diagnostics, therapeutics, and as research reagents.
XX
XX Example 15; Column 38; 32pp; English.
XX
XX The invention provides antisense compounds of 8-30 nucleotides that
XX inhibit the expression of human Cellular Inhibitor of Apoptosis-1
XX (c-IAP-1). The antisense compounds may be used for diagnostics,
XX therapeutics (for modulating the expression of c-IAP-1), prophylaxis
XX (e.g. to prevent or delay infection, inflammation, or tumor formation),
XX as research reagents (e.g. to distinguish between members of a
XX biological pathway) and in kits. Sequences AA22150-189 represent
XX phosphorothioate oligonucleotides used for antisense inhibition of
XX cellular inhibitor of apoptosis-1.
XX
XX Sequence 18 BP; 3 A; 2 C; 1 G; 12 T; 0 other;
XX
XX
XX Query Match 4.3%; Score 14; DB 20; Length 18;
XX Best Local Similarity 100.0%; Pred. No. 6.9e+03;
XX Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 160 aaataataataaa 173
XX |||||
XX Db 18 AAATATAAATAAA 5
XX
XX RESULT 18
XX AAS01590
XX ID AAS01590 standard; DNA; 22 BP.
XX
XX AAS01590;
XX
XX 18-JUL-2001 (first entry)
XX
XX Human IQGAP2 Cpg island 5'-bisulfite PCR primer.
XX
XX Human; T-type calcium channel; CACNA1G; cytosine methylation; Cpg island;
XX KM cellular proliferative disorder; colorectal cancer; age related disease;
XX KM apolipoprotein B; APOB; caudal type homeobox transcription factor 2;
XX KM CDX2; epidermal growth factor receptor; EGFR; fibrillin-1; FBN1;
XX KM G protein-coupled receptor 37; GPR37; heat shock 70KD protein 6; HSP70B;
XX KM HSPA6; RasGAP-related protein; IQGAP2; proteinase-activated receptor 2;
XX KM PAR2; paired-like homeodomain transcription factor 2; PITX2; Klotho; KL;
XX KM patched A; patched B; PTCHA; PTCHB; syndecan 1; syndecan 4; SDCL; SDCL4;
XX KM chromosome 5q; PCR primer; ss.
XX
XX Homo sapiens.
XX
XX OS
XX OS
XX WO200119845-A1.
```

XX 22-MAR-2001.
 PD 14-SEP-2000; 2000WO-US25479.
 XX 15-SEP-1999; 99US-0398522.
 XX (UYJO) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
 PA Issa J;
 PI WPI; 2001-244777/25.
 DR New nucleic acid molecule for use as a marker for screening cancer,
 PT comprises the coding region for a T-type calcium channel and regulatory
 PT sequences associated with the channel -
 PS Claim 21; Page 34; 125pp; English.
 XX The present sequence for 5'-bisulfite PCR primer is used to study the
 CC methylation state of human RasGAP-related protein (IQGAP2) which
 CC maps to chromosome 5q. The methylation state of specific regions
 CC within CpG islands associated with a novel T-type calcium channel
 CC CACNA1G gene correlate with several cancerous phenotypes involving
 CC various tissue and cell types. Since aberrant methylation of normally
 CC unmethylated CpG islands is often observed in immortalised and
 CC transformed cells, CACNA1G is implicated in cellular proliferative
 CC disorders e.g. leukaemia, colorectal, lung, breast and other cancers. The
 CC nucleic acid coding for CACNA1G is useful as a marker for screening
 CC cancer and age related diseases. A diagnostic kit containing primers
 CC (AAS01574-AAS01623) for amplification of a CpG-containing nucleic acid,
 CC where the primer hybridises with a target polynucleotide sequence
 CC (AAS01627-AAS01676), can be used for detecting aberrant methylation. The
 CC CpG island sequences (AAS01677-AAS01692) are selected from genes encoding
 CC CACNA1G, apolipoprotein B (APOB), caudal type homeobox transcription
 CC factor 2 (CDX2), epidermal growth factor receptor (EGFR), fibrillin-1
 CC (FBN1), G protein-coupled receptor 37 (GPR37), heat shock 70kD protein 6
 CC (HSP70B), HSPA6, RasGAP-related protein (IQGAP2), Klotho (KL),
 CC proteinase-activated receptor 2 (PAR2), paired-like homeodomain
 CC transcription factor 2 (PTRX2), patched A and B (PTCHA; PTCNB) and
 CC syndecan 1 and 4 (SDC1; SDC4) or a MINT31 sequence.
 CC Sequence 22 BP; 4 A; 0 C; 3 G; 15 T; 0 other;
 XX
 SO Query Match 4.3%; Score 14; DB 22; Length 22;
 Best Local Similarity 100.0%; Pred. No. 6.8e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209
 Db 2 ttattttatttta 15

RESULT 19
 AAS01643/c
 ID AAS01643 standard; DNA; 22 BP.
 XX AAS01643;
 AC
 XX 18-JUL-2001 (first entry)
 DT
 XX Human IQGAP2 5'-target sequence for bisulfite PCR.
 DE
 XX Human: T-type calcium channel; CACNA1G; cytosine methylation; CpG island;
 KW cellular proliferative disorder; colorectal cancer; age related disease;
 KW apolipoprotein B; APOB; caudal type homeobox transcription factor 2;
 KW CDX2; epidermal growth factor receptor; EGFR; fibrillin-1; FBN1;
 KW G protein-coupled receptor 37; GPR37; heat shock 70kD protein 6; HSP70B;
 KW HSPA6; RasGAP-related protein; IQGAP2; proteinase-activated receptor 2;
 KW PAR2; paired-like homeodomain transcription factor 2; PTRX2; Klotho; KL;
 KW patched A; patched B; PTCHA; PTCNB; syndecan 1; syndecan 4; SDC1; SDC4;
 KW chromosome 5q; ds.

XX Homo sapiens.
 OS
 XX W0200119645-A1.
 PN
 XX 22-MAR-2001.
 PD
 XX 14-SEP-2000; 2000WO-US25479.
 PF 15-SEP-1999; 99US-0398522.
 XX (UYJO) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
 PA Issa J;
 PI WPI; 2001-244777/25.
 DR New nucleic acid molecule for use as a marker for screening cancer,
 PT comprises the coding region for a T-type calcium channel and regulatory
 PT sequences associated with the channel -
 PS Claim 20; Page 37; 125pp; English.
 XX The present sequence for human RasGAP-related protein (IQGAP2)
 CC 5'-target sequence (complementary to the 5'-bisulfite PCR primer) is used
 CC to study the methylation state of IQGAP2 which maps to chromosome 5q. The
 CC methylation state of specific regions within CpG islands associated with
 CC a novel T-type calcium channel CACNA1G gene correlate with several
 CC cancerous phenotypes involving various tissue and cell types. Since
 CC aberrant methylation of normally unmethylated CpG islands is often
 CC observed in immortalised and transformed cells, CACNA1G is implicated in
 CC cellular proliferative disorders e.g. leukaemia, colorectal, lung, breast
 CC and other cancers. The nucleic acid coding for CACNA1G is useful as a
 CC marker for screening cancer and age related diseases. A diagnostic kit
 CC containing primers (AAS01574-AAS01623) for amplification of a
 CC CpG-containing nucleic acid, where the primer hybridises with a target
 CC polynucleotide sequence (AAS01627-AAS01676), can be used for detecting
 CC aberrant methylation. The CpG island sequences (AAS01677-AAS01692) are
 CC selected from genes encoding CACNA1G, apolipoprotein B (APOB), caudal
 CC type homeobox transcription factor 2 (CDX2), epidermal growth factor
 CC receptor (EGFR), fibrillin-1 (FBN1), G protein-coupled receptor 37
 CC (GPR37), heat shock 70kD protein 6 (HSP70B), HSPA6, IQGAP2, Klotho (KL),
 CC proteinase-activated receptor 2 (PAR2), paired-like homeodomain
 CC transcription factor 2 (PTRX2), patched A and B (PTCHA; PTCNB) and
 CC syndecan 1 and 4 (SDC1; SDC4) or a MINT31 sequence.
 CC Sequence 22 BP; 15 A; 3 C; 0 G; 4 T; 0 other;
 XX
 SO Query Match 4.3%; Score 14; DB 22; Length 22;
 Best Local Similarity 100.0%; Pred. No. 6.8e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209
 Db 21 ttattttatttttta 8

RESULT 20
 AAV07952/c
 ID AAV07952 standard; DNA; 26 BP.
 XX AAV07952;
 AC
 XX 02-FEB-1999 (first entry)
 DT
 XX Helicobacter pylori polypeptide GHP0 1414 5' DNA primer.
 DE
 XX GHP0 1414; infection; gastritis; ulcer; vaccine; diagnosis;
 KW therapy; PCR; primer; ss.
 KW Synthetic.
 OS Helicobacter pylori.

XX MO9843479-A1.
PN 08-OCT-1998.
PD
XX
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX WPI; 1998-568251/48.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 145; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07954) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07921) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73032) designated GHPO 1414.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter
CC infections.
XX
XX Sequence 26 BP; 14 A; 5 C; 4 G; 3 T; 0 other;
SQ

Query Match 4.3%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.7e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 307 ttltcatgtttct 320
Db 18 TTTTCATGTTTTCT 5

RESULT 21
AAV07922/C
ID AAV07922 standard; DNA; 26 BP.
XX
XX AAV07922;
AC
XX 02-FEB-1999 (first entry)
DT
XX
DE Helicobacter pylori polypeptide GHPO 386 5' DNA primer.
XX
KW GHPO 386; infection; gastritis; ulcer; vaccine; diagnosis; therapy;
KW PCR; primer; ss.
XX
XX Synthetic.
OS Helicobacter pylori.
OS
XX
XX WO9843479-A1.
PN
XX 08-OCT-1998.
PD
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX

DR WPI; 1998-568251/48.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 137; 184pp; English.
XX
XX This 5' primer is used with a 3' primer (see AAV07924) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV72001
CC) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73022) designated GHPO 386.
CC The isolated polynucleotide, and encoded polypeptide, can be used to
CC develop vaccines for the treatment and prevention of Helicobacter
CC infections.
XX
XX Sequence 26 BP; 15 A; 5 C; 4 G; 2 T; 0 other;
SQ

Query Match 4.3%; Score 14; DB 19; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.7e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 307 ttltcatgtttct 320
Db 20 TTTTCATGTTTTCT 7

RESULT 22
AAV07937/C
ID AAV07937 standard; DNA; 27 BP.
XX
XX AAV07937;
AC
XX 02-FEB-1999 (first entry)
DT
XX
DE Helicobacter pylori polypeptide GHPO 896 5' DNA primer.
XX
KW GHPO 896; infection; gastritis; ulcer; vaccine; diagnosis;
KW therapy; PCR; primer; ss.
XX
XX Synthetic.
OS Helicobacter pylori.
OS
XX
XX WO9843479-A1.
PN
XX 08-OCT-1998.
PD
XX 31-MAR-1998; 98WO-US06421.
PF
XX 01-APR-1997; 97US-0834666.
PR 01-APR-1997; 97US-0831310.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PI (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS ET VACCINS.
XX Al-Garawi A, Kleanthous H, Lissolo L, Miller C, Tomb J;
XX
XX WPI; 1998-568251/48.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastroduodenal diseases
XX
XX Claim 5; Page 141; 184pp; English.
XX
CC This 5' primer is used with a 3' primer (see AAV07939) in the PCR
CC amplification of Helicobacter, e.g. Helicobacter pylori, genomic
CC DNA in order to obtain DNA (see AAV07916) encoding the unprocessed
CC form of a 76 kDa polypeptide (see AAW73027) designated GHPO 896.
CC The isolated polynucleotide, and encoded polypeptide, can be used
CC to develop vaccines for the treatment and prevention of Helicobacter

CC infections.
 XX
 SO Sequence 27 BP; 14 A; 5 C; 4 G; 4 T; 0 other;

Query Match 4.3%; Score 14; DB 19; Length 27;
 Best Local Similarity 100.0%; Pred. No. 6.7e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 307 ttttcagtcttct 320
 Db 18 ttttcagtcttct 5

RESULT 23
 AAT42655/c
 ID AAT42655 standard; DNA; 29 BP.
 XX
 AC AAT42655;
 XX
 DT 25-FEB-1997 (first entry)
 XX
 DE Primer for amplifying verotoxin (VT-1) subunit A coding sequence.
 XX
 KW Verotoxin; Escherichia coli; enteric infection; diarrhoea; vaccine;
 KM haemolytic uraemic syndrome; detection; ss.
 XX
 OS Synthetic.
 XX
 PN M09630043-A1.
 XX
 PD 03-OCT-1996.
 XX
 PF 25-MAR-1996; 96MO-US04093.
 XX
 PR 24-MAR-1995; 95US-0410058.
 XX
 PA (OPHI-) OPHIDIAN PHARM INC.
 XX
 PI Carroll SB, Padhye NV, Stafford DC;
 PS
 DR WPI; 1996-505779/50.
 XX
 PT Compn. contg. neutralising antitoxin against E.coli vero-toxin -
 PT used to treat intoxicated individuals, and as a prophylactic against
 PT diarrhoeal disease or extra-intestinal complications of E.coli
 PT infection
 PS
 XX Example 6; Page 58; 101pp; English.
 CC Compositions containing neutralising antitoxin against one or more E.
 CC coli verotoxin (VT) can be used to treat intoxicated adults and
 CC children with enteric bacterial infections. They may also be used as
 CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the
 CC development of extra-intestinal complications of E.coli infection,
 CC especially haemolytic uraemic syndrome. The antitoxin can also be
 CC used to detect E. coli VT in a sample. The VT is recombinant,
 CC preferably a fusion protein containing a non-VT protein sequence and
 CC part of the E.coli VT1 or VT2 sequence. Two primers (AAT42655,
 CC AAT42656) were used to amplify the verotoxin VT-1 A subunit coding
 CC sequence and add a histidine tag coding sequence to the subunit
 CC sequence. Two primers (AAT42655, AAT42658) were used to amplify the
 CC verotoxin VT-1 A and B subunits and add a histidine tag coding
 CC sequence to the subunit sequences.
 XX
 SO Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.3%; Score 14; DB 17; Length 29;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 189 aaataattattt 202

Db 22 AAAAAAAAAA
 22 AAAAAAAAAA
 9

RESULT 24
 AAA51200/c
 ID AAA51200 standard; DNA; 29 BP.
 XX
 AC AAA51200;
 XX
 DT 26-SEP-2000 (first entry)
 XX
 DE N-terminal primer for E. coli verotoxin 1 subunit A gene.
 XX
 KW VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;
 KM recombinant production; screening; dairy; anti-bacterial; vaccine;
 KW primer; polynistidine; ss.
 XX
 OS Escherichia coli.
 OS Synthetic.
 OS
 PN US6080400-A.
 XX
 PD 27-JUN-2000.
 XX
 PF 13-MAR-1997; 97US-0816977.
 XX
 PR 24-MAR-1995; 95US-0410058.
 XX
 PA (OPHI-) OPHIDIAN PHARM INC.
 XX
 PI Williams JA, Byrne LM;
 XX
 DR WPI; 2000-451195/39.
 XX
 PT Bacterial cell for recombinantly expressing bacterial toxins in large
 PT quantities useful for immunization and treatment of bacterial
 PT infections, comprises expression vector encoding bacterial toxin
 PS
 XX Example 6; Column 83; 83pp; English.
 CC E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into
 CC pET-23b, designed to allow expression of the native proteins containing
 CC C-terminal polynistidine tags. The VT-1 and VT-2 genes were engineered
 CC to convert the signal sequence methionine codon into a NdeI site to
 CC allow cloning of the amplified genes into the vector without addition of
 CC vector-encoded amino acids. The C-terminal primers comprises the
 CC C-terminal 7 codons of each gene fused to the sequence CTCGAGCC, in order
 CC to add the polynistidine tag. The primers delete the native stop codons,
 CC and when cloned into pET-23 add a C-terminal extension of Leu-Glu-(His)6.
 CC VT B chains are small proteins (approximately 8 kDa), so use of a small
 CC affinity tag was preferred (i.e. polynistidine). A polynistidine affinity
 CC tag facilitates single step affinity purification of subunits from
 CC periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A
 CC and VT-2 A chains, expression of maltose binding protein (MBP) fused
 CC subunits was undertaken. Due to the toxicity of the VT-2 B subunit,
 CC strict uninduced promoter control is necessary to permit cell viability.
 CC Bacterial host cells expressing a recombinant expression vector encoding
 CC a polynistidine affinity tag and a portion of the VT-2 B chain are
 CC claimed. The vector is chosen from pET24hisVT2BL⁺, pET24hisVT2BL⁻ and
 CC pET24VT2, where "L⁺" indicates that the vector encodes the preprotein
 CC form of the protein and "L⁻" indicates that the vector encodes the mature
 CC form of the protein. The bacterial cell is capable of expressing large
 CC quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing
 CC non-mammals and for detecting bacterial toxins in environmental samples
 CC including soil, water, industrial samples, biological samples and samples
 CC obtained from food and dairy processing instruments.
 XX
 SO Sequence 29 BP; 11 A; 2 C; 5 G; 11 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 29;
 Best Local Similarity 100.0%; Pred. No. 6.6e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 189 aaataattattt 202
 |||||
 Db 22 AAAATAATTATTTT 9

RESULT 25

AAAT25703/C
 ID AAAT25703 standard: cDNA to mRNA: 31 BP.

AC AAAT25703;

DT 10-OCT-1996 (first entry)

DE Human gene signature HUNGSO7904.

KW Gene signature: messenger RNA; mRNA; relative abundance; frequency;
 human; cloning; mapping; non-biased library; diagnosis; detection;
 cell typing; abnormal cell function; ss.

OS Homo sapiens.

PN WO9514772-A1.

PD 01-JUN-1995.

PF 11-NOV-1994; 94WO-JP01916.

PR 12-NOV-1993; 93JP-0355504.

PA (MATS/) MATSUBARA K.

PA (OKUB/) OKUBO K.

PI Matsubara K, Okubo K;

DR WPI; 1995-206931/27.

PT Identifying gene signatures in 3'-directed human cDNA library - e.g.
 for diagnosis of abnormal cell function, by preparing cDNA that

PT reflects relative abundance of corresp. mRNA in specific human

PS tissues

CC Claim 1; Page 1910; 2245pp; Japanese.

CC A single-stranded DNA (or its complementary strand or the corresp.
 double-stranded DNA) which comprises one of the 7837 "GS" sequences

CC given in AAT19001-T26837 and which is able to hybridise to part of

CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)

CC sequences were obtained from 3'-directed cDNA libraries prepared

CC from various human tissues: synthesis of cDNA was initiated from the

CC 3'-end of mRNA by using poly(1) as the sole primer. Since the 3'-

CC untranslated sequence is unique to a particular mRNA species, almost

CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library

CC is constructed so as to reflect accurately the relative abundance of

CC different mRNAs in the particular tissue from which it was derived.

CC The appearance frequency of a given GS in a cDNA library can be

CC determined (esp. using primers and probes derived from the GS

CC sequences) as a means of diagnosing abnormal cell function or for

CC recognising different cell types.

CC Sequence 31 BP; 14 A; 3 C; 3 G; 10 T; 1 other;

CC Query Match 4.3%; Score 14; DB 16; Length 31;

CC Best Local Similarity 100.0%; Pred. No. 6.6e+03;

CC Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 198 attttattttaa 211
 |||||
 Db 20 ATTATTTATTTTAA 7

RESULT 26
 AAV67854
 ID AAV67854 standard: DNA: 31 BP.

AC AAV67854;

DT 24-DEC-1998 (first entry)

DE Nucleotide fragment containing polymorphic site, WI-11163.

XX ss; polymorphic site; nucleic acid analysis; diagnosis; monitoring;

XX cancer; inflammation; heart disease; CNS disease.

OS Homo sapiens.

PN WO9838846-A2.

PD 11-SEP-1998.

PF 06-MAR-1998; 98WO-US04571.

PR 28-MAR-1997; 97US-0042125.

PR 07-MAR-1997; 97US-0813159.

PA (AFRY-) AFFYMETRIX INC.

PI Berno A, Chee M, Fan J, Lipschutz RJ;

DR WPI; 1998-495419/42.

PT New nucleic acid segments containing polymorphic sites, or

PT complements and methods of detecting a nucleic acid - for general

PT use including diagnosis and monitoring of diseases

PS Claim 1; Page 25; 42pp; English.

CC New nucleic acid segment comprising one of the 10 - 100 bp sequences

CC given in the specification (sequences of a polymorphic site), or the

CC complement of the segment and a method of analysing a nucleic acid

CC comprising determining the base occupying the polymorphic site of the

CC polymorphic fragment sequences are disclosed in the specification. The

CC information obtained from nucleic acid analysis by the method described

CC is useful in diagnosis or monitoring of diseases like cancer,

CC inflammation, heart disease, CNS diseases, and susceptibility to

CC infection by microorganisms. In addition, the nucleic acid segments are

CC useful in manufacturing medication in the treatment of prophylaxis of

CC diseases, and also the use of the DNA segments as pharmaceutical.

CC Sequence 31 BP; 17 A; 2 C; 4 G; 7 T; 1 other;

CC Query Match 4.3%; Score 14; DB 19; Length 31;

CC Best Local Similarity 100.0%; Pred. No. 6.6e+03;

CC Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 209 aaattgtttaa 222
 |||||
 Db 17 aaattgtttaa 30

RESULT 27

AAZ27689/C
 ID AAZ27689 standard: DNA: 32 BP.

AC AAZ27689;

DT 22-DEC-1999 (first entry)

DE PCR primer for Verotoxin gene.

XX Verotoxin; VT1; VT2; detection; PCR primer; ss.

OS Synthetic.

```

OS  Escherichia coli.
XX
XX  JP11243996-A.
XX
XX  14-SEP-1999.
XX
XX  27-FEB-1998; 98UP-0047677.
XX
XX  27-FEB-1998; 98UP-0047677.
XX
XX  (TOYO ) TOYOBO KK.
XX
XX  WPI: 1999-603716/52.
XX
XX  An oligonucleotide for amplification of verotoxin - useful in the
XX  detection of inactivated verotoxin gene by transfer of a foreign DNA
XX  fragment
XX
XX  Claim 11; Page 9; 10pp; Japanese.
XX
XX  This sequence represents a PCR primer of the invention. The primer is
XX  used for amplification of the E. coli verotoxin (VT) gene. The
XX  oligonucleotide is useful for detection of inactivated VT gene by
XX  transfer of a foreign DNA fragment. Simple, rapid and specific
XX  amplification of VT gene from environmental factors is achieved using the
XX  oligonucleotide of the invention.
XX
XX  Sequence 32 BP; 12 A; 2 C; 4 G; 14 T; 0 other;
XX
XX  Query Match 4.3%; Score 14; DB 20; Length 32;
XX  Best Local Similarity 100.0%; Pred. No. 6.6e+03;
XX  Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0
XX
OY  189 aaataattatttt 202
XX  |||||||||||
DB  17 AAATAATTATTTT 4

RESULT 28
AAAT97603/C
ID AAAT97603 standard; DNA; 36 BP.
XX
XX  AAT97603;
AC
XX  30-APR-1998 (first entry)
XX
XX  Shigella dysenteriae delta-stx allele PCR primer 13.
XX
XX  Delta-virg allele; delta-guab-A allele; PCR; amplification; primer;
XX  delta-stx allele; shigellosis; vaccine; ss.
XX
XX  Synthetic.
OS  Shigella dysenteriae.
XX
XX  WO9737685-A1.
XX
XX  16-OCT-1997.
XX
XX  09-APR-1997; 97WO-US05954.
XX
XX  09-APR-1996; 96US-0629600.
XX
XX  (UYMA-) UNIV MARYLAND BALTIMORE.
XX
XX  Levine MM, Noriega FR;
XX
XX  WPI: 1997-512417/47.
XX
XX  Shigella mutants with mutation in guab-A - used in vaccines against
XX  Shigellosis
XX
XX  Example 6; Page 57; 94pp; English.
PS

```

Query Match	Best Local Similarity	Score	DB	Length	36;
Matches	14;	Conservative	0;	Mismatches	0;
				Indels	0;
				Gaps	0;
OY	189	aaataattattt	202		
DB	27	AAATAATTATTTT	14		
RESULT	29				
AAC90606	1D	AAC90606	standard; RNA; 36 BP.		
AC	AAC90606;				
XX	20-MAR-2001	(first entry)			
XX	Tomato spotted wilt virus S RNA partial sequence #10.				
XX	Tospovirus resistance; transgenic plant; tomato spotted wilt virus;				
XX	Impatiens necrotic spot virus; TSMV; ss.				
XX	Tomato spotted wilt virus.				
XX	OS				
XX	Key	Location/Qualifiers			
XX	misc_binding	1..13			
XX	misc_binding	/*tag= a			
XX	misc_binding	/bound_molecety= "binds nucleotides 33-21 of AAC89655"			
XX	misc_binding	15..28			
XX	misc_binding	/*tag= b			
XX	misc_binding	/bound_molecety= "binds nucleotides 19-6 of AAC89655"			
XX	misc_binding	32..36			
XX	misc_binding	/*tag= C			
XX	misc_binding	/bound_molecety= "binds nucleotides 5-1 of AAC89655"			
XX	US6150585-A.				
XX	21-NOV-2000.				
XX	26-NOV-1996;	96US-0757011.			
XX	02-MAY-1991;	91US-0694734.			
XX	14-APR-1993;	93US-0047346.			
XX	26-OCT-1993;	93US-0143397.			
XX	27-JUL-1994;	94US-0280903.			
XX	03-NOV-1994;	89US-0431259.			
XX	05-DEC-1989;	89US-0446024.			
XX	(NOVS.) NOVARTIS FINANCE CORP.				
XX	Peters D, Gielen JUL, De Haen PT, Van Grinsven MOUM, Kool AJ;				
XX	Goldbach RW;				
XX	WPI: 2001-060031/07.				
XX	Recombinant DNA construct comprising a DNA sequence encoding an RNA				
XX	sequence that codes for a tospovirus protein, useful for producing				
XX	plants with reduced susceptibility to tospovirus infection -				

XX Example 9; Fig 16c; 49pp; English.

CC The present invention provides DNA constructs encoding RNA sequences from
CC a tospovirus which can be used to produce transgenic plants with immunity
CC to tospoviruses. Examples of tospoviruses include the tomato spotted wilt
CC virus and the Impatiens necrotic spot virus.

SO Sequence 36 BP; 7 A; 0 C; 0 G; 29 U; 0 other;

Query Match 4.3%; Score 14; DB 22; Length 36;

Best Local Similarity 21.4%; Pred. No. 6.5e+03;
Matches 3; Conservative 11; Mismatches 0; Indels 0; Gaps 0;

OY 196 ttattttatttta 209

Db 7 uuuuuuuuuuuua 20

RESULT 30

AA062952

ID AA062952 standard; DNA; 37 BP.

XX

AA062952;

AC 09-SEP-1994 (first entry)

DE Glycophorin antibody 1C3 Fab coding region PCR primer.

XX Glycophorin; antibody 1C3; target binding polypeptide; PCR;

KW polymerase chain reaction; primer: antibody engineering;

KW humanized antibody; phagemid pHEA; plasmid p56;ss.

OS Synthetic.

PM WO9407921-A.

PD 14-APR-1994.

PF 24-SEP-1993; 93WO-AU00491.

PR 25-SEP-1992; 92AU-0004973.

PA (CSIR) COMMONWEALTH SCI & IND RES ORG.

PI Atwell JL, Colman PM, Hudson PJ, Irving RA, Kortt A;

PI Lah M, Malbyrl, Power BE;

DR WPI; 1994-135515/16.

XX New target-binding polypeptide(s) used for diagnosis, etc.

PT Having a stable core polypeptide region with at least one

PT target-binding region covalently attached, opt. mutated to alter

PT specificity, etc.

PS Example; Page 36; 67pp; English.

XX PCR primers given in AA062951-52 were used to clone anti-glycophorin

CC antibody 1C3 Fab coding region. The DNA sequence of the first 1443

CC bases of the Fab fragment in pHEA, ready for ligation post PCR

CC amplification for ligation into p569, is given in AA062958.

XX Sequence 37 BP; 14 A; 5 C; 0 G; 18 T; 0 other;

SO

RESULT 31

AAF55449/C

ID AAF55449 standard; DNA; 45 BP.

XX

AAF55449;

AC 29-MAY-2001 (first entry)

DE Oligonucleotide used to construct the shuttle vector pAAO-E-TATA.

XX Adenovirus vector; gene delivery vector; E2B gene; E4 gene; vaccine;

KW replication defective virus; cell proliferation; cell differentiation;

KW gene therapy; ss.

OS Synthetic.

PM EP1083229-A1.

PD 14-MAR-2001.

PF 10-SEP-1999; 99EP-0202966.

PR 10-SEP-1999; 99EP-0202966.

PA (INTR-) INTROGENE BV.

DR WPI; 2001-228258/24.

PT Producing a recombinant adenovirus-like gene delivery vehicle with

PT modified E2B or E4 functions, for gene therapy, comprises generating an

PT adenoviral vector where E2B or E4 is under the control of a synthetic

PT promoter

XX Example 1; Page 10; 56pp; English.

XX The specification describes a method for producing a recombinant

CC adenovirus-like gene delivery vehicle having reduced expression of

CC adenoviral E2B and/or E4 gene products in a target cell. The method

CC comprises generating a recombinant adenoviral vector lacking E1A and

CC E1B sequences, but having at least the E2B and/or E4 sequences encoding

CC products essential for adenoviral replication. Compared with previous

CC vectors, the new recombinant vectors are replication defective and

CC express the remaining viral genes only at background levels. The vector

CC itself does not dominantly elicit a response of the immune system, but

CC the immune response is directed primarily against the transgene product,

CC and is less toxic to the cells which, in turn, results in a prolonged

CC synthesis of the protein of interest. The recombinant adenoviral

CC vectors are used in the functional characterization of gene products in

CC cells, tissues or animals in order to find genes that encode for

CC proteins with a desired function such as those that interfere with cell

CC proliferation and differentiation. The vectors are further used as

CC vaccines, in gene therapy, and for protein production in mammalian

CC cells. Oligonucleotides AAF55449-50 were annealed together, and used to

XX construct a shuttle vector which was used in the method of the invention.

SO Sequence 45 BP; 14 A; 5 C; 7 G; 19 T; 0 other;

Query Match 4.3%; Score 14; DB 22; Length 45;

Best Local Similarity 100.0%; Pred. No. 6.4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 ctgaaattataa 92

Db 21 CTGAAATTATATAA 8

RESULT 32

AAF55450

ID AAF55450 standard; DNA; 45 BP.

CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.
XX
SQ Sequence 47 BP; 16 A; 4 C; 2 G; 25 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 6.4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 162 atataataaatt 175
|||||
Db 23 ATATATAATAAATT 10

RESULT 37
AAZ67533/C
ID AAZ67533 standard; DNA: 47 BP.
XX
AC AAZ67533;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:1880.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,C)
FT /*tag= a
/standard_name= "single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PE 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI; 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome -
XX
PS Claim 1; Page 631; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the

CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.
XX
SQ Sequence 47 BP; 15 A; 6 C; 6 G; 20 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 6.4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 95 atatgataagtata 108
|||||
Db 30 ATATGATATAAGTATA 17

RESULT 38
AAZ67549/C
ID AAZ67549 standard; DNA: 47 BP.
XX
AC AAZ67549;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human map-related biallelic marker SEQ ID NO:1896.
XX
KW Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT variation replace(24,G)
FT /*tag= a
/standard_name= "single nucleotide polymorphism"
XX
PN WO9954500-A2.
XX
PD 28-OCT-1999.
XX
PE 21-APR-1999; 99WO-IB00822.
XX
PR 21-APR-1998; 98US-0082614.
PR 23-NOV-1998; 98US-0109732.
XX
PA (GEST) GENSET.
XX
PI Cohen D, Blumenfeld M, Chumakov I;
XX
DR WPI; 2000-013267/01.
XX
PT Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome -
XX
PS Claim 1; Page 635; 2745pp; English.
XX
CC AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and

CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ. ID NOS 2857, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.
XX
XX Sequence 47 BP; 17 A; 10 C; 2 G; 18 T; 0 other;

	Query Match	4.38;	Score 14;	DB 21;	Length 47;
	Best Local Similarity	100.0%;	Pred. No. 6.4e+03;		
	Matches 14;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
OY	148 tttaaaaggaaa	161			
db	27 ttttAAAAAGGAAA	14			

RESULT	39	
AAZ67813		
ID	AAZ67813	standard; DNA: 47 BP.
XX		
AC		
XX	AAZ67813;	
DT	10-SEP-2001	(first entry)
XX		
DE		
XX		
XX	Human map-related diallelic marker	SEQ ID NO:2160.
KW	Human genome; diallelic marker; high density disequilibrium map;	
KW	genomic map; haplotype; phenotype; polymorphic base; genotyping;	
KW	haplotyping; hybridisation; identification; characterisation;	
KW	diagnosis; single nucleotide polymorphism; SNP; ds.	
XX		
OS	Homo sapiens.	
XX		
FH	Key	Location/Qualifiers
FT	Variation	replac(24,6)
FT		/tag= a
FT		/standard_name= "single nucleotide polymorphism"
XX		
PN	W09954500-A2.	
XX		
PD	28-OCT-1999.	
XX		
PF	21-APR-1999;	99WO-IB00822.
XX		
PR	21-APR-1998;	98US-0082614.
PR	23-NOV-1998;	98US-0109732.
XX		
PA	(GEST) GENSET.	
XX		
PI	Cohen D, Blumenfeld M, Chumakov I;	
XX		
DR	WPI: 2000-013267/01.	
XX		
PT	Novel diallelic markers used to construct a high density disequilibrium	
PT	map of the human genome	-
XX		
PS	Claim 1; Page 691; 2745pp; English.	
XX		
CC	AAZ65554 to AAZ69578 represent human diallelic markers from the present	
CC	invention, which contain a polymorphic base at position 24 of their	
CC	nucleotide sequences. AAZ69579 to AAZ77440 represent amplification	
CC	primers for the diallelic markers. The diallelic markers of the	
CC	invention have a variety of uses: they can be used for high density	
CC	mapping of the human genome, and in complex association studies and	
CC	haplotyping studies which are useful in determining the genetic basis	
CC	for disease states. Compositions and methods of the invention can also	
CC		

CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.
XX
S0 Sequence 47 BP; 19 A; 4 C; 8 G; 16 T; 0 other;

```

Query Match          4.3%; Score 14; DB 21; Length 47;
Best Local Similarity 100.0%; Pred. No. 6 4e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 198 attttattttaa 211
    |||
Db 11 attttattttaa 24

```

RESULT 40
 ID AAA98312
 AC AAA98312 standard; DNA; 50 BP.
 XX AAA98312;
 XX 02-FEB-2001 (first entry)
 DE Human MSH6 fragment 8/exon 8 to 10 DNA Ref-Seq fragment.
 XX
 KW Human mismatch repair gene; hMSH6; disease predisposition; genotype;
 KW mutation; carcinoma; colorectal; endometrial; ovarian; leukemia;
 KW neoplastic disease; drug development; ds.
 XX
 OS Homo sapiens.
 PN DEJ909878-AI.
 XX
 PD 07-SEP-2000.
 XX
 PF 06-MAR-1999; 99DE-1009878.
 XX
 PR 06-MAR-1999; 99DE-1009878.
 XX
 PA (UYDR) UNIV DRESDEN TECH.
 XX
 PI Plaschke J, Kruppa C, Schackert H;
 DR WPI: 2000-588378/56.
 XX
 PT Novel variants of the human mismatch repair gene, MSH6, useful e.g. for
 PT determining predisposition to cancer and for development of drugs
 PS
 PS Disclosure: Fig 3; 14pp; German.

CC development of pharmaceuticals: (ix) developing diagnostic kits and other
CC systems for genotyping; and (x) developing in vivo and in vitro test
CC systems for expressing individual forms of the MSH6 gene, e.g. for
CC studying pathophysiology of disease or processes in which MSH6 is
CC involved, and for drug development and testing.

XX Sequence 50 BP; 8 A; 8 C; 5 G; 29 T; 0 other;

Query Match 4.3%; Score 14; DB 21; Length 50;

Best Local Similarity 100.0%; Pred. No. 6.3e+03;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 140 cttttgttttaa 153

11 cttttgttttaa 24

Db

RESULT 41

AAFA8097 standard; DNA; 15 BP.

AAFA8097;

30-MAR-2001 (first entry)

IGFBP3 oligonucleotide #1517.

Antisense therapy: antiproliferative; antiinflammatory; antipsoriatic;
cytostatic; dermatological; cardiant; virucide; ophthalmological; keloid;
skin disorder; insulin-like growth factor 1 receptor; IGF-1; pityriasis;
IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris;
growth factor mediated cell proliferation; ichthyosis; serborrhea; ruba;
keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease;
hyperneovascular condition; hyperplasia; kidney disease;
neovascular condition of the retina; ss.

Homo sapiens.

MO200078341-A1.

28-DEC-2000.

21-JUN-2000; 2000MO-AU00693.

21-JUN-1999; 99US-0140345.

(MURD-) MURDOCH CHILDRENS RES INST.

Wright CJ, Werther GA, Edmondson SR;

WPI; 2001-041421/05.

Ameliorating the effects of a disorder, e.g. psoriasis, by
administering UV (ultra-violet) treatment (optional) and an antisense
nucleic acid that inhibits or reduces growth factor mediated cell
proliferation and/or inflammation -

Example 7; Page 54; 201pp; English.

CC The present invention relates to a method for ameliorating the effects
CC of skin disorders. The method comprises contacting the skin with an
CC antisense oligonucleotide, (for Insulin-like Growth Factor [IGF]-1
CC receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of
CC inhibiting or reducing growth factor mediated cell proliferation,
CC inflammation and/or other disorders. The present sequence is an
CC oligonucleotide which can be used to design the antisense
CC oligonucleotides of the present invention (see AAF45151 and
CC AAF45153-F45161). The method is useful for ameliorating the effects of
CC psoriasis, ichthyosis, pityriasis, ruba, pilaris, serborrhea, keloids,
CC keratosis, neoplasias, scleroderma, warts, benign growths, cancers of the
CC skin, a hyperneovascular condition such as a neovascular condition of the
CC retina, brain or skin, growth factor-mediated malignancies, other

CC sclerotic disease, kidney disease, hyperproliferation of the inside of
CC blood vessels or any other hyperplasia.

XX Sequence 15 BP; 7 A; 1 C; 3 G; 4 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 agaagaatgttta 38

3 agaagaatgttta 15

Db

RESULT 42

AAFA8098 standard; DNA; 15 BP.

AAFA8098;

30-MAR-2001 (first entry)

IGFBP3 oligonucleotide #1518.

Antisense therapy: antiproliferative; antiinflammatory; antipsoriatic;
cytostatic; dermatological; cardiant; virucide; ophthalmological; keloid;
skin disorder; insulin-like growth factor 1 receptor; IGF-1; pityriasis;
IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris;
growth factor mediated cell proliferation; ichthyosis; serborrhea; ruba;
keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease;
hyperneovascular condition; hyperplasia; kidney disease;
neovascular condition of the retina; ss.

Homo sapiens.

MO200078341-A1.

28-DEC-2000.

21-JUN-2000; 2000MO-AU00693.

21-JUN-1999; 99US-0140345.

(MURD-) MURDOCH CHILDRENS RES INST.

Wright CJ, Werther GA, Edmondson SR;

WPI; 2001-041421/05.

Ameliorating the effects of a disorder, e.g. psoriasis, by
administering UV (ultra-violet) treatment (optional) and an antisense
nucleic acid that inhibits or reduces growth factor mediated cell
proliferation and/or inflammation -

Example 7; Page 54; 201pp; English.

CC The present invention relates to a method for ameliorating the effects
CC of skin disorders. The method comprises contacting the skin with an
CC antisense oligonucleotide, (for Insulin-like Growth Factor [IGF]-1
CC receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of
CC inhibiting or reducing growth factor mediated cell proliferation,
CC inflammation and/or other disorders. The present sequence is an
CC oligonucleotide which can be used to design the antisense
CC oligonucleotides of the present invention (see AAF45151 and
CC AAF45153-F45161). The method is useful for ameliorating the effects of
CC psoriasis, ichthyosis, pityriasis, ruba, pilaris, serborrhea, keloids,
CC keratosis, neoplasias, scleroderma, warts, benign growths, cancers of the
CC skin, a hyperneovascular condition such as a neovascular condition of the
CC retina, brain or skin, growth factor-mediated malignancies, other
CC blood vessels or any other hyperplasia.

Sequence 15 BP; 7 A; 0 C; 3 G; 5 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 agaagaatgttta 38

|||||

2 agaagaatgttta 14

Db

RESULT 43

AAFA8099

AAFA8099 standard; DNA; 15 BP.

AC AAF48099;

30-MAR-2001 (first entry)

IGFBP3 oligonucleotide #1519.

Antisense therapy; antiproliferative; antiinflammatory; antipsoriatic; cytosolic; dermatological; cardiant; virucide; ophthalmological; keloid; skin disorder; insulin-like Growth Factor 1 receptor; IGF-1; ptyriasis; IGF binding protein; IGFBP-2; IGFBP3; inflammation; psoriasis; pilaris; growth factor mediated cell proliferation; ichthyosis; serborrhoea; ruba; keratosis; neoplasia; scleroderma; wart; skin cancer; sclerotic disease; hypervascular condition; hyperplasia; kidney disease; neovascular condition of the retina; ss.

Homo sapiens.

WO200078341-A1.

28-DEC-2000.

21-JUN-2000; 2000WO-AU00693.

21-JUN-1999; 99US-0140345.

(MORD-) MURDOCH CHILDRENS RES INST.

Wraight CJ, Werther GA, Edmondson SR;

WPI: 2001-041421/05.

Ameliorating the effects of a disorder, e.g. psoriasis, by administering UV (ultra-violet) treatment (optional) and an antisense nucleic acid that inhibits or reduces growth factor mediated cell proliferation and/or inflammation -

Example 7; Page 54; 201pp; English.

The present invention relates to a method for ameliorating the effects of skin disorders. The method comprises contacting the skin with an antisense oligonucleotide, (for insulin-like Growth Factor [IGF]-1 receptor, IGF binding protein [IGFBP]-2 or IGFBP3), which is capable of inhibiting or reducing growth factor mediated cell proliferation, inflammation and/or other disorders. The present sequence is an oligonucleotide which can be used to design the antisense oligonucleotides of the present invention (see AAF45151 and AAF45153-45161). The method is useful for ameliorating the effects of psoriasis, ichthyosis, ptyriasis, ruba, pilaris, serborrhoea, keloids, keratosis, neoplasia, scleroderma, warts, benign growths, cancers of the skin, a hypervascular condition such as a neovascular condition of the retina, brain or skin, growth factor-mediated malignancies, other sclerotic disease, kidney disease, hyperproliferation of the inside of blood vessels or any other hyperplasia.

Sequence 15 BP; 6 A; 0 C; 4 G; 5 T; 0 other;

Query Match 4.0%; Score 13; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 agaagaatgttta 38

|||||

1 agaagaatgttta 13

Db

RESULT 44

AAA57758/C

AAA57758 standard; DNA; 16 BP.

AC AAA57758;

20-OCT-2000 (first entry)

Nucleotide sequence which is bound by 22 domain of RIP60 polypeptide.

Human; RIP60; zinc finger protein; nucleic acid delivery complex;

nucleic acid binding domain; nucleic acid condensation domain; ss.

Synthetic.

WO200040723-A2.

13-JUL-2000.

04-JAN-2000; 2000WO-US00212.

04-JAN-1999; 99US-0114743.

04-JAN-1999; 99US-0114745.

(UYVE-) UNIV VERMONT & STATE AGRIC COLLEGE.

Heintz NH, Houchens CR;

WPI: 2000-465985/40.

Non-viral nucleic acid delivery complex for delivering a nucleic acid molecule into a cell comprises a modular polypeptide -

Example 17; Page 74; 115pp; English.

AAA57752-66 represent sequences which are bound by the 22 domain of the human RIP60 polypeptide. RIP60 is a zinc finger protein. The nucleic acid binding domain of the RIP60 polypeptide is used to construct a non-viral nucleic acid delivery complex comprising a modular polypeptide. The complex comprises a modular peptide containing a nucleic acid binding domain and a nucleic acid condensation domain that bind with and condense a nucleic acid molecule of more than 50 kilobases in length. The complex also comprises one or more polypeptides selected from a cell recognition domain, a protein transduction domain, a protein degradation domain, an intracellular targeting domain, a protein interaction domain, an epitope domain and a protein purification domain. The complexes are used to deliver a nucleic acid to a cell. The nucleic acids delivered are of various sizes and preferably greater than 50 kilobases, especially more than 100 or more than 200 kilobases in length.

Sequence 16 BP; 5 A; 0 C; 0 G; 11 T; 0 other;

Query Match 4.0%; Score 13; DB 21; Length 16;

Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 110 ttaataataataa 122

|||||

15 TTAATTAATAATAA 3

Db

RESULT 45

```

AA092084/c
ID  AA092084 standard; cDNA; 17 bp.
XX
AC  AA092084;
XX
DT  07-JAN-1996 (first entry)
XX
DE  Renilla reniformis luciferase DNA probe-1.
XX
KW  Luciferase; enzyme; bioluminescence; luminescence; label; DNA probe;
KW  antibody; oligonucleotide; ss.
XX
OS  Synthetic.
XX
PN  US5418155-A.
PD  23-MAY-1995.
XX
PF  29-DEC-1989; 89US-0458952.
XX
PR  29-DEC-1989; 89US-0458952.
PR  20-AUG-1992; 92US-0933017.
PR  17-JUN-1993; 93US-0079700.
PR  14-DEC-1993; 93US-0167650.
XX
PA  (UYGE-) UNIV GEORGIA RES FOUND INC.
XX
PI  Cormier MJ, Lorenz WN;
XX
DR  WPI; 1995-199740/26.
XX
PT  New recombinant Renilla luciferase polypeptide - used as a
PT  luminescent tag, partic in bio-luminescence assays and for the prodn
PT  of antibodies
XX
PS  Disclosure; Fig. 4; 18bp; English.
XX
CC  This 17-mer oligonucleotide DNA probe, along with Probe-2 (AA092085)
CC  are used to screen an R. reniformis cDNA library to isolate cDNA
CC  encoding Renilla luciferase. The luciferase was then expressed
CC  using E. coli.
XX
SQ  Sequence 17 bp; 6 A; 0 C; 2 G; 9 T; 0 other;

Query Match          4.0%; Score 13; DB 16; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  271 aaaaattatctt 283
    |||||||
DB  15 AAAAAATTATTT 3

```

Search completed: January 24, 2002, 03:28:19
 Job time: 3671 sec

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:22:17 : Search time 216.42 Seconds
 (without alignments)
 356.526 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
 Sequence: 1 atgaaaaaatatttccaa.....gtccaatgaagaagatgca 90

Scoring table: IDENTITY_NUC
 Gapop 10.0, Gapext 1.0

Searched: 930621 seqs, 428662619 residues

Total number of hits satisfying chosen parameters: 1661242

Minimum DB seq length: 0
 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Listing first 45 summaries

Database :

N.Geneseq_1101.*
 1: /SIDS2/gcgdata/geneseq/NA1980.DAT.*
 2: /SIDS2/gcgdata/geneseq/NA1981.DAT.*
 3: /SIDS2/gcgdata/geneseq/NA1982.DAT.*
 4: /SIDS2/gcgdata/geneseq/NA1983.DAT.*
 5: /SIDS2/gcgdata/geneseq/NA1984.DAT.*
 6: /SIDS2/gcgdata/geneseq/NA1985.DAT.*
 7: /SIDS2/gcgdata/geneseq/NA1986.DAT.*
 8: /SIDS2/gcgdata/geneseq/NA1987.DAT.*
 9: /SIDS2/gcgdata/geneseq/NA1988.DAT.*
 10: /SIDS2/gcgdata/geneseq/NA1989.DAT.*
 11: /SIDS2/gcgdata/geneseq/NA1990.DAT.*
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 13: /SIDS2/gcgdata/geneseq/NA1992.DAT.*
 14: /SIDS2/gcgdata/geneseq/NA1993.DAT.*
 15: /SIDS2/gcgdata/geneseq/NA1994.DAT.*
 16: /SIDS2/gcgdata/geneseq/NA1995.DAT.*
 17: /SIDS2/gcgdata/geneseq/NA1996.DAT.*
 18: /SIDS2/gcgdata/geneseq/NA1997.DAT.*
 19: /SIDS2/gcgdata/geneseq/NA1998.DAT.*
 20: /SIDS2/gcgdata/geneseq/NA1999.DAT.*
 21: /SIDS2/gcgdata/geneseq/NA2000.DAT.*
 22: /SIDS2/gcgdata/geneseq/NA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	90	100.0	1392	20	AAK60299
2	60	66.7	327	20	AAK60300
3	32.6	36.2	9542	20	AAK20260
4	32.6	36.0	4015	22	AAK01490
5	31.6	35.1	6250	21	AAK62306
6	31.2	34.7	3337	17	AAK34620
7	31.2	34.7	3337	20	AAK15174
8	31.2	34.7	3337	22	AAH76457
9	30.8	34.2	13830	21	AAK35351
10	30.6	34.0	936	22	AAK02659
11	30.6	34.0	936	22	AAK58252

C 12	30.6	34.0	936	22	AAK58254	Oligonucleotide D1
C 13	30.6	34.0	936	22	AAK58257	Oligonucleotide D1
C 14	30.6	34.0	936	22	AAK58259	Oligonucleotide D2
C 15	30.6	34.0	936	22	AAK58262	Oligonucleotide D2
C 16	30.6	34.0	936	22	AAK58255	Oligonucleotide D1
C 17	30.6	34.0	936	22	AAK58255	Oligonucleotide D1
C 18	30.6	34.0	26928	18	AAK33003	Human prothrombin
C 19	30.6	34.0	87350	18	AAK33003	Human prothrombin
C 20	30.4	33.8	1347	20	AAK13328	Enterococcus faeca
C 21	30.4	33.8	2364	21	AAK70246	Plasmodium falcipa
C 22	30.3	33.3	244	20	AAK0301	Fragment of the be
C 23	30.3	33.3	244	22	AAK58238	Oligonucleotide D1
C 24	30.3	33.3	4140	18	AAK77331	Solanum tuberosum
C 25	29.8	33.1	210	20	AAK61492	B. burgdorferi ant
C 26	29.8	33.1	324	20	AAK61491	B. burgdorferi ant
C 27	29.8	33.1	784	22	AAH05038	Human cDNA clone (
C 28	29.8	33.1	3454	22	AAH18466	Human cDNA sequenc
C 29	29.8	33.1	14417	22	AAI62923	Human genomic DNA
C 30	29.8	33.1	14426	22	AAI62921	Human genomic DNA
C 31	29.6	32.9	513445	22	AAK1373	Soybean 318013 reg
C 32	29.6	32.9	1518	17	AAK37313	Aromatic acyl tran
C 33	29.6	32.9	1592	21	AAK27085	Human secreted pro
C 34	29.6	32.9	1703	12	AAQ15131	Clone pAC1 encodi
C 35	29.6	32.9	1703	19	AAV15701	Zucchini ACC synth
C 36	29.6	32.9	1703	22	AAK04541	Zucchini 1-aminocy
C 37	29.6	32.9	5994	21	AAK23618	Zucchini ACC synth
C 38	29.6	32.9	110000	22	AAK70222	Plasmodium falcipa
C 39	29.6	32.9	393	21	AAK84800	Nucleotide sequenc
C 40	29.4	32.7	546	22	AAK09430	Human secreted pro
C 41	29.4	32.7	1219	21	AAK48516	Human cDNA clone (
C 42	29.4	32.7	1222	21	AAK45059	Arabidopsis thalia
C 43	29.4	32.7	1223	21	AAK40150	Arabidopsis thalia
C 44	29.4	32.7	1405	22	AAH34185	Human colon cancer
C 45	29.4	32.7	1978	22	AAH14169	Human cDNA sequenc

ALIGNMENTS

RESULT 1
 AAK60299 Standard; DNA; 1392 BP.
 ID AAK60299
 AC AAK60299;
 DT 12-AUG-1999 (first entry)
 DE DNA encoding the beta-2 toxin of Clostridium perfringens type C.
 KW Beta-2 toxin; Clostridium perfringens type C; gene promoter;
 KW vaccine; Clostridium tetani; ss.
 OS Clostridium perfringens.
 PN FR2768747-A1.
 PD 26-MAR-1999.
 PF 19-SEP-1997; 97FR-0011710.
 PR 19-SEP-1997; 97FR-0011710.
 PA (INSP) INST PASTEUR.
 PI Gilbert M, Popoff MR;
 DR WPI: 1999-217498/19.
 DR P-PSDB: AAK16591.
 PT Clostridium beta2 toxin gene promoter and signal sequence - usefu
 PT against toxins from Clostridium perfringens
 PS Example A: Page 31; 46pp; French.

RESULT	3
DB	268 atgtaaaataattctcaagttctacgtaatttttagttcttcacatgtttctctatcgtc 337
AAx20260/C	
ID	AAx20260 standard; DNA; 9542 BP.
XX	
AC	AAx20260;
XX	

DT 04-May-1999 (first entry)
 XX
 DE Borrelia burgdorferi polynucleotide sequence #13.
 XX
 KN Borrelia burgdorferi, spirochete; bacterium; pathogen; Lyme disease.

kw	infection; diagnosis; characterisation; detection; ds.
xx	
os	<i>Borrelia burgdorferi</i> .
vv	

PN	W09858943-A1.
XX	
PD	30-DEC-1998.

PF	18-JUN-1998;	98WC-US12764
XX	03-SEP-1997;	97US-0057483
PR	20-JUN-1997;	97US-0050359
PR	12-SEP-1997;	97US-0050359

PR	22-JUL-1997;	97US-0053377
XX		
PA	(HUMA-) HUMAN GENOME SCI IN	

XX Clayton R, Dougherty BA, Fraser C, Iathigra R, Smith HO,
PI White OR;
PI
XX

XX	PT	PI
New isolated <i>Borrelia burgdorferi</i> nucleic acids - used to develop products for the detection, diagnosis, characterisation, prevention		

PS Claim 1, Page 920-925; 1128pp; English.

CC Bb infections, e.g. Lyme disease. They can also be used for the
CC production of biosynthetic products, e.g. enzymes. *Borrelia* belongs
CC to a family of motile, spiral-shaped bacteria called Spirochetes.

CC endemic relapsing fever, and Lyme borreliosis, more commonly known as

XX Sequence 9542 BP; 3812 A; 1160 C; 1113 G; 3457 T; 0 other;
SQ

Best Local Similarity 60.9%; Pred. NO. 22;
Matches 53; Conservative 0; Mismatches 34; Indels 0; Gaps
QY 4 aaaaattatctcaagttactgtaatttcatgtttcctaattgtga 63
||||| || |

Db 3872 GAAC TATTATTAATAAAAAAGGAGAGCA 3846

RESULT	4
AAS01490/c	
ID	AAS01490 standard; DNA; 4015 BP.

XX	AA01490;
AC	
XX	18-JUL-2001 (first entry)
DT	
XX	
DE	Human secreted protein gene #31.
XX	
KW	Human secreted protein; gene therapy; autoimmune disease;
KW	hyperproliferative disorder; cardiovascular disorder;
KW	cerebrovascular disorder; nervous system disorder; infection;
KW	ocular disorder; wound healing; epithelial cell proliferation;
KW	skin aging; transplantation; tissue regeneration; chemotaxis;
KW	food additive; preservative; ds.
OS	
XX	Homo sapiens.
PN	MO200123402-AI.
PD	
XX	05-APR-2001.
PF	
XX	26-SEP-2000; 2000WO-US26376.
PR	
XX	27-SEP-1999; 99US-0155808.
XX	
PA	(HUMA-) HUMAN GENOME SCI INC.
P1	
XX	Rosen CA, Ruben SM, Komatsoulis GA;
DR	
XX	WPI: 2001-266138/27.
DR	
P-PSDB:	AAM01070.
PT	
XX	Nucleic acids encoding 43 human secreted polypeptides, useful for
PT	preventing, diagnosing and/or treating e.g. cancers, Parkinson's
PT	disease and diabetic retinopathy -
XX	
PS	Claim 4; Page 452-453; 516pp; English.
XX	
CC	AA01460-AA01502 encode for novel human secreted proteins. The
CC	invention relates to 43 novel human secreted proteins (AAM01040-AAM01082)
CC	and their gene sequences which can be used in gene therapy. The secreted
CC	proteins are useful to prevent, treat or ameliorate a medical condition
CC	in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or
CC	sheep. The secreted proteins are also useful in diagnosing a pathological
CC	condition or susceptibility to a pathological condition. Antibodies to
CC	the secreted proteins can also be used in alleviating symptoms associated
CC	with disorders and in diagnostic immunoassays e.g. radioluminoassays or
CC	enzyme linked immunosorbent assays (ELISA). Disorders which are diagnosed
CC	or treated include autoimmune diseases e.g. rheumatoid arthritis,
CC	hyperproliferative disorders e.g. neoplasms of the breast or liver,
CC	cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders
CC	e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g.
CC	Parkinson's disease, infections caused by bacteria, viruses and fungi and
CC	ocular disorders e.g. corneal infection. The polypeptides can also be
CC	used to aid wound healing and epithelial cell proliferation, to prevent
CC	skin aging due to sunburn, to maintain organs before transplantation, for
CC	supporting cell culture of primary tissues, to regenerate tissues and in
CC	chemotaxis. The polypeptides can also be used as a food additive or
CC	preservative to increase or decrease storage capabilities.
XX	
SO	Sequence 4015 BP; 1122 A; 712 C; 743 G; 1438 T; 0 other;
Query Match	36.0%; Score 32.4; DB 22; Length 4015;
Best Local Similarity	68.2%; Pred. No. 24;
Matches	45; Conservative 0; Mismatches 21; Indels 0; Gaps 0
OY	2 tgaataaaatttcacagtttactgtaatttatgtttcattcatgtttctatgttg 61
Dd	3088 TGAGGAAGAAGTTATGAAAAGTTCACAGTCATTGTATCTTCATTTCATGTTAGTG 3029
OY	62 gagcaa 67
Dd	3028 GAGCTA 3023

Query Match	35.1%	Score 31.6	DB 21	Length 6250
Best local Similarity	62.8%	Pred. No. 36		
Matches 49	Conservative 0	Mismatches 29	Indels 0	Gaps 0
QY 4	aaaaaaatattccaagttctgtaatttaagtttcaagtttctattgtgga	63		
DB 853	AAATAAATTTTGAAATAATCTACAAAGTTTTTTTTTTCATTTCCTGGAATAGTT	794		
QY 64	gcaataagtcacatgaaa	81		
DB 793	GTAATAATGTGAAAAAAA	776		

```

ID   AAT34620 standard; DNA; 3337 BP.
XX
XX   AAT34620;
AC
XX
XX   12-NOV-1996 (first entry)
DT
XX
XX   P. vivax ESP-1 blood stage antigen coding sequence.
DE
XX
XX   ESP-1; blood stage antigen; diagnosis; malaria; infection;
KM   causative agent; antibody; monoclonal; polyclonal; assay; ds.
XX
XX   Plasmodium vivax (clone PvMB3.3.1).
OS
XX
XX   Key      Location/Qualifiers
FH   Exon     1..91
FT   Exon     /tag= a
FT   Exon     /note= "encodes initial (N-terminal) sequence of
FT   Intron    92..230
FT   Intron    /tag= b
FT   Intron    /note= "contains typical malaria intervening
FT   Exon     231..3197
FT   Exon     /tag= c
XX
XX   US5532133-A.
XX
XX   02-JUL-1996.
XX
XX   02-JUN-1993; 9305-0072610.
XX
XX   02-JUN-1993; 9305-0072610.
XX
XX   (UNYNY ) UNIV NEW YORK STATE.
XX
XX   Barnwell JW;
XX
XX   WPI; 1996-321110/32.
XX
XX   P-PSDB; AAR98747.
XX
XX   Antibodies to Plasmodium vivax blood stage antigens - used to
PT   diagnose malaria and to determine whether P. vivax is the species
PT   responsible for infection
XX
XX   Example 4; Column 15-20; 22pp; English.
XX
XX   The present sequence encodes a species-specific Plasmodium vivax
CC   malarial antigen, PvESP-1. The gene appears to be missing a small
CC   portion of its 5' end. This protein is secreted into the plasma of
CC   a susceptible mammalian host after infection. Monoclonal/polyclonal
CC   antibodies can be utilized in assays used to diagnose malaria, as well
CC   as to determine whether P. vivax is the species responsible for the
CC   infection.
XX
XX   Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
SQ
XX
XX   Query Match      34.7%; Score 31.2; DB 17; Length 3337;
XX   Best Local Similarity 63.2%; Pred. No. 44;
XX   Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
XX
XX   0y   10 atattcaaaagttacttaatttattggtttcatggtttctattgttgagcaata 69
XX   Db   138 AATATATATAGTATATATATACAGATTATTTATTTATCTTACTGTTGACATGA 79
XX
XX   0y   70 agtccaatgaagaacaa 85
XX   Db   78 ATTAAAGAAAGCAAA 63
XX
XX   RESULT 7
XX   AAX15174/c
XX   ID AAX15174 standard; DNA; 3337 BP.

```

```

XX
XX   AAX15174;
AC
XX
XX   28-APR-1999 (first entry)
DT
XX
XX   DNA encoding a secreted blood-stage protein called PvESP-1.
DE
XX
XX   Erythrocyte secreted protein-1; PvESP-1; malarial antigen;
KM   blood-stage protein; malaria; monoclonal antibody 1D11G10; ds.
XX
XX   Plasmodium vivax.
OS
XX
XX   Key      Location/Qualifiers
FH   CDS      1..3197
FT   CDS      /tag= a
FT   CDS      /note= "contains 1 intron"
FT   Exon     1..91
FT   Exon     /tag= b
FT   Exon     /number= 1
FT   Intron    92..230
FT   Intron    /tag= c
FT   Exon     /number= 1
FT   Exon     231..3194
FT   Exon     /tag= d
XX
XX   US5874527-A.
XX
XX   23-FEB-1999.
XX
XX   30-SEP-1996; 96US-0719822.
XX
XX   02-JUN-1993; 93US-0072610.
XX
XX   07-JUN-1995; 95US-0478417.
XX
XX   30-SEP-1996; 96US-0719822.
XX
XX   (UNYNY ) UNIV NEW YORK STATE.
XX
XX   Barnwell JW;
XX
XX   WPI; 1999-180063/15.
XX
XX   P-PSDB; AAW97039.
XX
XX   Plasmodium vivax peptide antigen - for diagnosis of malaria caused
PT   by Plasmodium vivax
XX
XX   Example 4; Fig 5A-C; 23pp; English.
XX
XX   The present sequence encodes a C-terminal erythrocyte secreted
CC   protein-1 (PvESP-1) of Plasmodium vivax. PvESP-1 is a malarial
CC   antigen which is a secreted blood-stage protein present in detectable
CC   amounts in biological samples from individuals infected with P. vivax.
CC   The protein comprises an epitope not present in other Plasmodium species
CC   that cause malaria in humans, and is bound by monoclonal antibody
CC   1D11G10. The peptide antigen can be used in immunoassays for diagnosis
CC   of malaria caused by P. vivax and/or can be used to produce antibodies
CC   for use in such immunoassays.
XX
XX   Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
SQ
XX
XX   Query Match      34.7%; Score 31.2; DB 20; Length 3337;
XX   Best Local Similarity 63.2%; Pred. No. 44;
XX   Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
XX
XX   0y   10 atattcaaaagttacttaatttattggtttcatggtttctattgttgagcaata 69
XX   Db   138 AATATATATAGTATATATATACAGATTATTTATTTATCTTACTGTTGACATGA 79
XX
XX   0y   70 agtccaatgaagaacaa 85
XX   Db   78 ATTAAAGAAAGCAAA 63
XX
XX   RESULT 7
XX   AAX15174 standard; DNA; 3337 BP.

```



```
RESULT 8
AAH76457/c
ID AAH76457 standard; DNA; 3337 BP.
XX
AC AAH76457;
XX
DT 22-OCT-2001 (first entry)
XX
DE Plasmodium vivax ESP-1 DNA.
XX
KW Plasmodium vivax: ESP-1; erythrocyte secreted protein-1; PvESP-1;
KW species-specific; malarial peptide antigen; infection; diagnosis;
KW malaria; ds.
XX
OS Plasmodium vivax.
XX
FH Key Location/Qualifiers
FT CDS 1..3197
FT /tag= a
FT /product= "ESP-1"
FT exon 1..91
FT /tag= b
FT /number= 1
FT intron 92..230
FT /tag= c
FT /number= 1
FT exon 231..3197
FT /tag= d
FT /number= 2
XX
XX US6231861-B1.
XX
PD 15-MAY-2001.
XX
PF 05-JUN-1998; 98US-0092458.
XX
PR 02-JUN-1993; 93US-0072610.
PR 07-JUN-1995; 95US-0478417.
PR 30-SEP-1996; 96US-0719822.
XX
PA (UYNV ) UNIV NEW YORK STATE.
XX
XX Barnwell JW;
XX
DR WPI: 2001-335068/35.
DR P-PSDB: AAG66528.
XX
XX New species-specific Plasmodium vivax malarial peptide antigens,
XX proteins or fragments secreted into the plasma of susceptible mammalian
XX host after infection, useful for diagnosing malaria
XX
XX Example 4; Fig 5; 23p; English.
XX
XX The invention relates to novel species-specific Plasmodium vivax
XX malarial peptide antigens which are proteins or fragments of
XX proteins secreted into the plasma of a susceptible mammalian host after
XX infection, and to monoclonal or polyclonal antibodies directed against
XX those antigens. The peptide antigens, monoclonal antibodies, and/or
XX polyclonal antibodies are useful in assays to diagnose malaria, and to
XX determine which P. vivax species is responsible for the infection.
XX The present sequence encodes P. vivax erythrocyte secreted
XX protein-1 (PvESP-1), a secreted species-specific blood stage
XX antigen provided in the invention.
XX
XX Sequence 3337 BP; 1304 A; 467 C; 875 G; 691 T; 0 other;
XX
Query Match 34.7%; Score 31.2; DB 22; Length 3337;
Best Local Similarity 63.2%; Pred. No. 44;
Matches 48; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
OY 10 attattcaagttactgtaatttattgtttcattgtttctattgttgagcacaata 69
```

```
DB 138 AATATATATGTTATATATAGCACTTTTATAGATTATTTTATCTTACTGTGAACGATCA 79
OY 70 agtccaatgaaagca 85
DB 78 ATTAAAGAAAGCA 63
RESULT 9
AAC35351
ID AAC35351 standard; DNA; 1226 BP.
XX
AC AAC35351;
XX
DT 17-OCT-2000 (first entry)
XX
DE Arabidopsis thaliana DNA fragment SEQ ID NO: 9873.
XX
KW Hybridisation assay; genetic mapping; gene expression control;
KW protein identification; signal transduction pathway;
KW metabolic pathway; promoter; termination sequence; ss.
XX
XX Arabidopsis thaliana.
XX
XX EP1033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
XX 25-FEB-1999; 99US-0121825.
XX 05-MAR-1999; 99US-0123180.
XX 09-MAR-1999; 99US-0123548.
XX 23-MAR-1999; 99US-0125788.
XX 25-MAR-1999; 99US-0126264.
XX 29-MAR-1999; 99US-0126785.
XX 01-APR-1999; 99US-0127462.
XX 06-APR-1999; 99US-0128234.
XX 08-APR-1999; 99US-0128714.
XX 16-APR-1999; 99US-0129845.
XX 19-APR-1999; 99US-0130077.
XX 21-APR-1999; 99US-0130449.
XX 23-APR-1999; 99US-0130510.
XX 23-APR-1999; 99US-0130891.
XX 28-APR-1999; 99US-0131449.
XX 30-APR-1999; 99US-0132048.
XX 30-APR-1999; 99US-0132407.
XX 04-MAY-1999; 99US-0132485.
XX 05-MAY-1999; 99US-0132485.
XX 06-MAY-1999; 99US-0132486.
XX 06-MAY-1999; 99US-0132487.
XX 07-MAY-1999; 99US-0132863.
XX 11-MAY-1999; 99US-0134256.
XX 14-MAY-1999; 99US-0134218.
XX 14-MAY-1999; 99US-0134221.
XX 14-MAY-1999; 99US-0134370.
XX 18-MAY-1999; 99US-0134768.
XX 19-MAY-1999; 99US-0134941.
XX 20-MAY-1999; 99US-0135124.
XX 21-MAY-1999; 99US-0135353.
XX 24-MAY-1999; 99US-0135629.
XX 25-MAY-1999; 99US-0136021.
XX 27-MAY-1999; 99US-0136392.
XX 28-MAY-1999; 99US-0136782.
XX 01-JUN-1999; 99US-0137222.
XX 03-JUN-1999; 99US-0137528.
XX 04-JUN-1999; 99US-0137502.
XX 07-JUN-1999; 99US-0137724.
XX 08-JUN-1999; 99US-0138094.
XX 10-JUN-1999; 99US-0138540.
XX 10-JUN-1999; 99US-0138847.
XX 14-JUN-1999; 99US-0139119.
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PR 16-JUN-1999; 990S-0139452.
PR 16-JUN-1999; 990S-0139453.
PR 17-JUN-1999; 990S-0139492.
PR 18-JUN-1999; 990S-0139454.
PR 18-JUN-1999; 990S-0139455.
PR 18-JUN-1999; 990S-0139456.
PR 18-JUN-1999; 990S-0139457.
PR 18-JUN-1999; 990S-0139458.
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PR 26-OCT-1999; 990S-0161361.
PR 28-OCT-1999; 990S-0161920.
PR 28-OCT-1999; 990S-0161992.
PR 28-OCT-1999; 990S-0161993.
PR 29-OCT-1999; 990S-0162142.

Query Match 34.2%; Score 30.8; DB 21; Length 1226;
Best Local Similarity 70.7%; Pred. No. 53;
Matches 41; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

OY 29 taattttatgatttcacgatttcttattgttgagcaataagtcacatgaagcaag 86.
Db 1157 ttatttttttttcacgatttctcgttaattgaagcagtaataataacagcgag 1214

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RESULT 10
AAD02659/C
ID AAD02659 standard; DNA; 13830 BP.
XX
AC AAD02659;
XX
DT 02-MAY-2001 (first entry)
XX
DE Tomato chromosome 5 harbouring the RIN and MC genes.
XX
KM Tomato; RIN; ripening inhibitor; MC; macrocalyx; sepal development;
XX senescence; pathogen infection; ethylene response; transgenic plant; ds.
XX
OS Lycopersicon esculentum.
XX
FH Key
FT misc_signal
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FT /note= "Putative transcription start site of RIN"
FT 1..5694
FT /tag= b
FT /note= "Transcribed region of RIN gene"
FT 211..5489
FT /tag= c
FT /product= "Tomato ripening-inhibitor (RIN) protein"
FT /note= "The specification states that the RIN gene
FT has 9 exons and 8 introns, however the sequence
FT represented in the figure 7 shows 10 exons and
FT 9 introns"
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FT /tag= d
FT /number= "1"
FT 396..3268
FT /tag= e
FT /number= "1"
FT 3269..3347
FT /tag= f
FT /number= "2"
FT 3348..3834
FT /tag= g
FT /number= "2"
FT 3835..3897
FT /tag= h
FT /number= "3"
FT 3898..3919
FT /tag= i
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FT 3920..3923
FT /tag= j
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FT 3924..4149
FT /tag= k
FT /number= "4"
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FT /tag= l
FT /number= "5"
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FT 4532..4572
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FT /tag= r
FT /number= "8"
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FT /number= "8"
FT 4863..4992
FT /tag= t
FT /number= "9"
FT 4993..5404
FT /tag= u
FT /number= "9"
FT /note= "The rin mutation begins at a point within
FT this region"
FT 5405..5489
FT /tag= v
FT /number= "10"
FT 5695..8237
FT /tag= w
FT /note= "MC promoter sequence; This region separates
FT the RIN from the MC transcribed region"
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FT /tag= x
FT /note= "Putative transcription start site of MC"
FT 8238..13830
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FT /note= "Transcribed region of MC gene"
FT 8251..13552
FT /tag= z
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FT /note= "The coding region has 8 exons and is interrupted
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FT /tag= aa
FT /number= "1"
FT 8440..10594
FT /tag= ab
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FT /note= "The rin mutation terminates within this
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FT /tag= ad
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FT /tag= ae
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FT /tag= ag
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FT /tag= aj
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FT 12560..12600
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FT 12601..12879
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FT /number= "6"
FT 12880..12992
FT /tag= am
FT /number= "7"
FT 12993..13449
FT intron

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PT a single surface -
XX
PS Example 6; Page 127; 159pp; English.
XX
CC The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
CC monitoring gene expression.
XX
SQ Sequence 936 BP; 4 A; 144 C; 7 G; 5 T; 776 other;

Query Match	34.0%;	Score 30.6;	DB 22;	Length 936;
Best Local Similarity	3.7%;	Pred. No. 59;		
Matches	3;	Conservative	62;	Mismatches 16;
				Indels 0;
				Gaps 0;

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OY      1 atgaaaaaatattctcaagttaacgtatcttttgcatttcataccttg   60  
Db      456 mmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmm    397
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OY      61 ggaagcataagtcocatgaaa       81  
Db      396 TAACGAmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmm    376
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Query Match	34.0%;	Score 30.6;	DB 22;	Length 936;
Best Local Similarity	3.7%;	Pred. No. 59;		
Matches	3;	Conservative	62;	Mismatches 16;
				Indels 0;
				Gaps 0.

Oy	1 atgaaaaaatatttccaagttaacgtatcttttttgcttcgaagttccatcgtc 60
	: :
Db	456 www 397
Oy	61 ggaagcataagtccaatgaaa 81
	: :
Db	396 TTAAGCWWW 376

Query Match	34.0%;	Score 30.6;	DB 22;	Length 936;
Best Local Similarity	3.7%;	Pred. No. 59;		
Matches	3;	Conservative	62;	Mismatches 16;
				Indels 0;
				Gaps 0

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Oy      1 atgaaaaaatatctcaagtttaccgtatttttgatttcacgtttccatcgtc 60
        : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      456 wwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwwww 397
Oy      61 ggagcataagtccaatgaaa 81
        |||:::~: ~::~~
Db      396 TAAGCWWWWWWWWWWWWW 376
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RESULT 15

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AAF58262/C
ID AAF58262 standard; DNA; 936 BP.
XX
AC AAF58262;
XX
DT 24-APR-2001 (first entry)
XX
DE Oligonucleotide D2007.
XX
KM Electron-transfer group; ETM; mismatch; genotyping;
XX gene expression; ss.
XX
OS Synthetic.
XX
PN WO200107665-A2.
XX
PD 01-FEB-2001.
XX
PF 26-JUL-2000; 2000WO-US20476.
XX
XX 26-JUL-1999; 99US-0145695.
PR 17-MAR-2000; 2000US-0190259.
XX
XX (CLIN-) CLINICAL MICRO SENSORS INC.
XX
XX UmeK RM;
XX
DR WPI; 2001-159728/16.
XX
PT Nucleic acids containing electron-transfer group, useful as labels in
PT hybridization assays, e.g. for genotyping, allowing repeat analyses on
PT a single surface -
XX
PS Example 6; Page 128; 159pp; English.
XX
CC The present invention relates to a composition comprising two nucleic
CC acids each containing an electron-transfer group (ETM) having
CC different redox potentials. The invention is used for electronic
CC detection of nucleic acids, especially of substitutions (mismatches)
CC and single-nucleotide polymorphisms, e.g. for genotyping,
CC monitoring gene expression.
XX
SQ Sequence 936 BP; 5 A; 139 C; 10 G; 6 T; 776 other;

Query Match 34.0%; Score 30.6; DB 22; Length 936;
Best Local Similarity 3.7%; Pred. No. 59;
Matches 3; Conservative 62; Mismatches 16; Indels 0; Gaps 0;

Oy 1 atgaaaaaatattcaagttactgaatttattgtttcattgttcttattgtt 60
   ::::::::::::::::::::: : : : : : : : : : : : : : : : : : :
Db 456 wmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmmm 397
   |||::: : : : : :
Oy 61 ggagcaataagtcacatgaaa 81
   |||::: : : : : :
Db 396 TAAGCWMWMMWMMWMMWMMWMMWMMWMMWMMWMMWMMWMMWMMWMMWMM 376

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Search completed: January 24, 2002, 02:22:23
Job time: 1891 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:24:45 ; Search time 1502.85 Seconds
(without alignments)
987.954 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaataatttcaaa.....gtccatgaagcaagtga 90

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1472140 seqs, 8248589755 residues

Total number of hits satisfying chosen parameters: 2944280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: gb_ba: *
2: gb_hcg: *
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11: gb_sts: *
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13: gb_un: *
14: gb_vi: *
15: em_ba: *
16: em_fun: *
17: em_hum: *
18: em_in: *
19: em_om: *
20: em_ov: *
21: em_ov: *
22: em_pat: *
23: em_ph: *
24: em_pl: *
25: em_ro: *
26: em_sts: *
27: em_sy: *
28: em_un: *
29: em_vi: *
30: em_htgo_hum: *
31: em_htgo_inv: *
32: em_htgo_rod: *
33: em_htg_hum: *
34: em_htg_inv: *
35: em_htg_rod: *
36: em_htg_other: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	90	100.0	90	6	AX004615
2	90	100.0	1392	1	L77965
3	90	100.0	1392	6	AX004613
4	88.4	98.2	54310	1	AP003515
5	60	66.7	327	6	AX004614
6	37.8	42.0	128806	9	AC021648
7	36.4	40.4	152816	2	AC079739
8	36.4	40.4	165176	2	AC022199
9	36.4	40.4	167322	2	AC016788
10	36.4	40.4	189772	2	AP003462
11	36.4	40.4	193415	2	AC074180
12	35.4	39.3	148	8	CNS01BGA
13	35.4	39.3	33460	8	AF099924
14	35.4	39.3	69431	3	AF067612
15	35.4	39.3	132319	9	AC006364
16	35.4	39.3	172766	2	AC079401
17	35.4	39.3	276829	2	AC006741
18	35.4	39.3	299864	2	AC006702
19	35	38.9	139927	9	AL354985
20	34.8	38.7	134478	8	AP000423
21	34.6	38.4	16005	3	AF049132
22	34.6	38.4	142805	2	AC022655
23	34.6	38.4	176633	2	AC017017
24	34.6	38.2	138781	9	AL157821
25	34.4	38.2	135410	2	AC016459
26	34.2	38.0	185591	2	AC016878
27	34.2	38.0	185702	2	AP002963
28	34.2	38.0	188517	2	AP003973
29	34.2	38.0	200875	2	AL591711
30	34.2	38.0	225016	2	AC063867
31	34.2	38.0	269661	2	AC023911
32	34.2	37.8	176343	9	CNS01DX3
33	34	37.6	9091	3	AF324831
34	33.8	37.6	204759	2	AC013285
35	33.8	37.6	245802	2	AC006279
36	33.6	37.3	193520	2	AL589787
37	33.6	37.3	204652	2	PFMAL13P6
38	33.6	37.1	2108	3	PF627
39	33.4	37.1	8597	2	PF627
40	33.4	37.1	88695	2	AC017727
41	33.4	37.1	120863	3	AC092399
42	33.4	37.1	163359	2	AC025699
43	33.4	37.1	175765	2	AC021111
44	33.4	37.1	193873	2	AC009775
45	33.4	37.1	193873	2	AC009775

ALIGNMENTS

RESULT	1	AX004615	90 bp	DNA	PAT	24-AUG-2000
LOCUS	AX004615	Sequence 3 from Patent WO9915669.				
DEFINITION	AX004615					
ACCESSION	AX004615.1	GI:9928056				
VERSION						
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
FEATURES						
source						

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Matches 90; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaatttcaagtttactgtaattttatglttcttcttctt 60
    |||||||
DB 1 ATGAAAAAATTTATTTCAAGTTTACTGTAATTTATGTTTCATGTTTCTTATTTGTT 60

QY 61 ggagcaataagtcacatgaaagcaagtga 90
    |||||||
DB 61 GGAGCAATAAGTCCAAATGAAGCAAGTGA 90

RESULT 2
LOCUS      L77965      1392 bp      DNA      BCT      28-JUL-1998
DEFINITION Clostridium perfringens C beta 2 toxin gene, complete cds.
ACCESSION      L77965
VERSION      L77965.1 GI:3342214
KEYWORDS
SOURCE
ORGANISM      Clostridium perfringens C.
               Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
               Clostridium.
REFERENCE      1 (bases 1 to 1392)
AUTHORS      Gilbert,M., Jolivet-Reynaud,C. and Popoff,M.R.
TITLE      Beta2 toxin, a novel toxin produced by Clostridium perfringens
JOURNAL      Gene 203 (1), 65-73 (1997)
MEDLINE      98085977
REMARK      Erratum: [[published erratum appears in Gene 1998 Mar
               27;210(1):173]]
REFERENCE      2 (bases 1 to 1392)
AUTHORS      Popoff,M.R.
TITLE      Direct Submission
JOURNAL      Submitted (15-JAN-1998) Toxines Microbiennes, Institut Pasteur,
               Paris cedex 15 75724, France
               GDSB:S:76036.
COMMENT      [Flatfile retrieved from GDSB Fri Jul 24 15:39:17 1998].
               Location/Qualifiers
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DB 268 ATGAAAAAATTTATTTCAAGTTTACTGTAATTTATGTTTCTTATTTGTT 327

QY 61 ggagcaataagtcacatgaaagcaagtga 90
    |||||||
DB 328 GGAGCAATAAGTCCAAATGAAGCAAGTGA 357

RESULT 3
LOCUS      AX004613      1392 bp      DNA      PAT      24-AUG-2000
DEFINITION Sequence 1 from Patent WO9915669.
ACCESSION      AX004613
VERSION      AX004613.1 GI:9928053
KEYWORDS
SOURCE      Clostridium perfringens.
ORGANISM      Clostridium perfringens
               Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
               Clostridium.
REFERENCE      1 (bases 1 to 1392)
AUTHORS      Gilbert,M. and Popoff,M.R.
TITLE      Clostridium toxin, and method for preparing immunogenic
               compositions
JOURNAL      Patent: WO 9915669-A 1 01-APR-1999;
               GIBERT MARYSE (FR); PASTEUR INSTITUT (FR)
FEATURES
      source
      1. .1392
         /organism="Clostridium perfringens"
         /db_xref="taxon:1502"
         268..1065
         /note="unnamed protein product"
         /codon_start=1
         /transl_table=11
         /protein_id="CAC04901.1"
         /db_xref="GI:9928054"
         /translation="MKIIISKFTVIFMFSCFLIVGALISPMKASAKETDAYRKVMENYL
         NALKNYDINTVNISEDERVNNVEOYREMLDFKYPNOOLKSEFILNSOKSDNKEIF
         NVKTEFLNGAIYDMEFTYSSKDKLIVSDMERTKVENGGKYLTPSPFTQVCTMDDEL
         AQAIGGVYPQYSDRFTYADNIIILNFQVATSGSRDLKVEYSVDHMMKDDVKASQ
         MVTGONPDSARQIRLITKGSQSFYKIRIRINFTPASIRVGEYGCA"
BASE COUNT      606 a          115 c          209 g          462 t
ORIGIN

Query Match      100.0%; Score 90; DB 6; Length 1392;
Best Local Similarity 100.0%; Pred. No. 1.9e-10;
Matches 90; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaatttcaagtttactgtaattttatglttcttcttctt 60
    |||||||
DB 268 ATGAAAAAATTTATTTCAAGTTTACTGTAATTTATGTTTCTTATTTGTT 327

QY 61 ggagcaataagtcacatgaaagcaagtga 90
    |||||||
DB 328 GGAGCAATAAGTCCAAATGAAGCAAGTGA 357

RESULT 4
LOCUS      AP003515      54310 bp      DNA      Circular      BCT      10-AUG-2001
DEFINITION Clostridium perfringens plasmid pcpl3 DNA, complete sequence.
```


ACCESSION AP003515 GI:15076712
VERSION
KEYWORDS Clostridium perfringens (strain:13) plasmid:pcp13 DNA.
SOURCE Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae; Clostridium.
ORGANISM 1 (bases 1 to 54310)
REFERENCE Ohnari,K., Ohshima,S., Hirakawa,H., Ohshima,K., Shiba,T., Shimizu,T., Hattori,M., Kuhara,S., Hayashi,H. and Shimizu,T.
AUTHORS Complete Nucleotide Sequence of the Virulence Plasmid pcp13 from Clostridium perfringens
TITLE Unpublished
JOURNAL 2 (bases 1 to 54310)
REFERENCE Shimizu,T.
AUTHORS Direct Submission
TITLE Submitted (12-APR-2001) Tohru Shimizu, Institute of Basic Medical Sciences, University of Tsukuba, Department of Microbiology; 1-1-1 Tennohda, Tsukuba, Ibaraki 305-8575, Japan
JOURNAL (E-mail:tshimizu@md.tsukuba.ac.jp, Tel:81-298-53-3354, Fax:81-298-53-3354)
AUTHORS
TITLE
FEATURES
SOURCE 1. Location/Qualifiers
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/plasmid="pcp13"
/strain="13"
/db_xref="taxon:1502"
/note="anaerobic pathogen for gas gangrene"
940.1692
/gene="soj"
940.1692
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/note="250 aa, similar to pir:140444 SpooA activation inhibitor soj from Bacillus subtilis (253 aa); 37% identity in 250 aa overlap
PCP01
para family"
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/transl_table=1
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/protein_id="BAB62438.1"
/db_xref="GI:15076713"
/translation="MKKISVENIKGVAKTSTANFGACLEKGRVLYVDLPSNLTKLEKAYMEDVSTADVLKNDLKEVTKTDPENDILPSNTLAPAFKTLDDNRSQONRIAKALEEIEDKIDCLIDCPALNKITYNALCASDEVLPPIKIDFALDGLILDSIEIKDEENPNLNFKGCFTMDSTTVNKVIOELKSVLGEKMFNTSIHQNIK VESTFECEPVFSSKARASLNKYKDLSEIF"
1751.3031
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1751.3031
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/note="426 aa, similar to gnu:AF300944_3 presumptive ParB protein from Lactococcus lactis subsplactis (242 aa); 30% identity in 266 aa overlap
PCP02"
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/transl_table=1
/product="ParB protein"
/protein_id="BAB62439.1"
/db_xref="GI:15076714"
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3147.3509
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3147.3509
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/note="120 aa, no significant homology"

/codon_start=1
/transl_table=1
/product="hypothetical protein"
/protein_id="BAB62440.1"
/db_xref="GI:15076715"
/translation="MEKTLAEKRINISPKKNGALVTTLYLPPKMLEVIGITENERECFYEIEDAKIKSEKQSEFAKEKTIISPKSTFTYLNKKLELYLGVSEDRSCITIEL RKDITLVKNDGRILDI"
3773.4024
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3773.4024
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/note="83 aa, similar to pir:R14710 probable transposase from Yersinia pestis (402 aa); 44% identity in 50 aa overlap
truncated"
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/db_xref="GI:15076717"
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4040.4222
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4040.4222
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/note="60 aa, similar to gp:AF143819.1 transposase-1 like protein from Escherichia coli (402 aa); 38% identity in 60 aa overlap
truncated"
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4588.4746
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4588.4746
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/translation="MLYTVNIESLNSQFRKFTYKLIFFNDVSLIKMLYLATEKVNK KWRNYPN"
complement(5169.5804)
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complement(5169.5804)
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/function="ATP-binding protein"
/note="211 aa, similar to gp:AP001508.4 ABC transporter (ATP-binding protein) from Bacillus halodurans (213 aa); 49% identity in 214 aa overlap"
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/transl_table=1
/product="probable ABC transporter"
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/db_xref="GI:15076720"
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complement(5804.7966)
/gene="PCP08"
complement(5804.7966)
CDS

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/gene="PCP08"
/Note="720 aa, similar to gp:Ap001508.3 BH0280 gene
product from Bacillus halodurans (713 aa); 23% identity in
661 aa overlap"
/codon_start=1
/transl_table=11
/product="conserved hypothetical protein"
/protein_id="BAB62446.1"
/db_xref="GI:15076721"
/translation="MKKVALILFIIITLISFEGYVSRNTEEMKLNOCNLENE
EYIISPDRENTNIAVESTIKSLDENGFSEVDINEDVKYINIGFSKRENEH
IPVSGREFHEKNDISYLSITDSBDTQDQIVINDENGNHEIRITKSLINFEEN
LEIYOENEOILDKLIEDLKSESIYOKRVMSQYNETKILLVFCILIMFIE
YOVLSYKKGIOKLLGHSSTFYMKEKLELVRIEIVLVLTLLVFNKFNPSLF
WKEMELICIVSIMITFIIVSVIIPYIVYSKITLSNIKKRPKSIITINSIVAIL
ASITLIFPSNADDSISCKGEKKNKWEERKOYIIPELGPNDSIOSPEIEMEK
EBAVILYRNKOCAILADENRTEPTSMEEKOMLPEEMRETIIVPNILKRVIVD
GNINISEDEKRIILVPEKYNFEKEILEYGYNSQESCTTSHKADKLN,VE
OKOKIIMKSNOKYPSYLDVNEPEGNYVTDIVSVLESNDKLSYKIIIGYNSPF
KIRANSEEVINGLEKYYDMYVLIIDPYLVYNASTIINIKAKYKVIIFAILVILAV
ISIILONTSLYENONKNKIYKLGHYRLIYRNMVFIMVLITWCPLAIASLITKD
INIIYFTLIVLYELVFIFENINSLEKNLIVINIGY"
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/db_xref="GI:15076722"
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L"
8779.9012
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8779.9012
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/Note="77 aa, similar to probable transposase from
Yersinia pestis plasmid pMT1 (402 aa); 25% identity in 158
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/product="probable transposase"
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/db_xref="GI:15076723"
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MKNLLEKREKWSKYPNVYKSWKMDMNLSTFF"
9127.9366
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Query Match      98.2%; Score 88.4; DB 1; Length 54310;
Best Local Similarity 98.9%; Pred. No. 3.9e-10;
Matches 89; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 atgaaaaaataattctcaagttactgaatttatgtttcattcagtttcttattgtt 60
|||||
DB 13654 ATGAAAAAATATTTTCAAGTTTACTGTAATTTTATGTTTTCATATTTTCTATTGTT 13713
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QY 61 ggagcaataagtcacatgaagaagcaagtgc 90
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DB 13714 GGAGCAATAAGTCACATGAAGAAGCAAGTCA 13743
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RESULT 5
AX004614 327 bp DNA PAT 24-AUG-2000
LOCUS AX004614
DEFINITION Sequence 2 from Patent WO9915669.
ACCESSION AX004614
VERSION AX004614.1 GI:9928055
KEYWORDS

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SOURCE
ORGANISM
Clostridium perfringens.
Clostridium perfringens
Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
Clostridium.
REFERENCE
1 (bases 1 to 327)
AUTHORS
Gibert,M. and Popoff,M.R.
TITLE
Clostridium toxin, and method for preparing immunogenic
compositions
JOURNAL
Patent: WO 9915669-A 2 01-APR-1999;
GIBERT MARYSE (FR); PASTEUR INSTITUT (FR)
FEATURES
source
1.327
/organism="Clostridium perfringens"
/db_xref="taxon:1502"
BASE COUNT
141 a 13 c 44 g 129 t
ORIGIN

Query Match      66.7%; Score 60; DB 6; Length 327;
Best Local Similarity 100.0%; Pred. No. 0.00053;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atgaaaaaataattctcaagttactgaatttatgtttcattcagtttcttattgtt 60
|||||
DB 268 ATGAAAAAATATTTTCAAGTTTACTGTAATTTTATGTTTTCATGTTTCTATTGTT 327
|||||

RESULT 6
AC021648/c
LOCUS AC021648/c
DEFINITION
Human BAC Library) complete sequence.
ACCESSION AC021648
VERSION AC021648.25 GI:13899174
KEYWORDS
HTG.
SOURCE
ORGANISM
human.
Human.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 128809)
AUTHORS
Muzny,D.M., Adams,C., Adio-Odola,B., Ali-Osman,F.R., Allen,C.,
Aldbrooks,S.L., Amaratunga,H.C., Are,J.R., Banks,T., Barbara,J.,
Benton,J., Blumage,K., Blankenburg,K., Bonnin,D., Bouck,J.,
Bowle,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,
Burich,P., Burkett,C., Burrell,K.L., Byrd,N.C., Caron,T.F.,
Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,R.,
Chen,Z., Chiu,D., Chowdhury,I., Christopoulos,C., Cleveland,C.D.,
Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,
Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,
Denu,A.L., Ding,Y., Dinh,H.H., Douhaite,K.J., Draper,H.,
Dugan-Rocha,S., Durbin,K.J., Earhart,C., Edgar,D., Edwards,C.C.,
Elhaj,C., Emerling,S., Escotto,M., Falls,T., Ferraguto,D.,
Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J.,
Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guenara,W.,
Gunnarane,P., Hale,S., Hamilton,K., Han,J., Harris,C., Harris,K.,
Hart,M., Haylak,P., Hawes,A., Hernandez,J., Hernandez,B.,
Hodgson,A., Hognes,M., Holloway,C., Hollins,P., Homsi,F.,
Howard,S., Huber,J., Huily,S., Hume,J., Ioshikhes,I., Jackson,L.E.,
Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S.,
Karlsson,E., Kelly,S., Khan,U., King,L., Kovari,J., Kovar,C.,
Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lee,E., Lewis,L.C.,
Lewis,L., Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W.,
Lounseged,H., Lozano,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,
Ma,J., Maheshwari,M., Mapua,P., Marondei,I., Martin,R.,
Martindale,A., Martinez,E., Massey,E., Mawhney,E., McLeod,M.P.,
Meador,M., Mei,G., Merscher,S., Metzger,M., Miller,K., Miner,G.,
Miner,Z., Mitchell,T., Mohabbat,K., Montgomery,K.T., Morgan,N.,
Morris,S., Moser,M., Neal,D., Nelson,D., Newton,J., Newton,M.,
Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokenko,S.,
Ogunh,M., Okunolu,G., Orangunye,N., Oviedo,R., Pace,A., Payton,B.,
Peery,J., Perez,U., Peters,L., Pickens,R., Primus,E., Pu,L.L.,
Quiles,M., Ren,Y., Rivers,M., Rojss,A., RojudoKan,I., Rolfe,M.,
Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shim,C.,

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Shooshart, N., Sisson, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Swalek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, Y., Villalon, D., Vinson, R., Wall, R., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleceyk, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y. F., Zhou, J., Zorilla, S., Zorichlepati, R., and Gibbs, R.

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
AUTHORS
TITLE
JOURNAL
COMMENT

Unpublished
2 (bases 1 to 128809)
Worley, K.C.
Direct Submission
Submitted (19-JAN-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 128809)
Worley, K.C.
Direct Submission
Submitted (01-MAY-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On May 1, 2001 this sequence version replaced gi:13876397.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as low coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: http://gc.bcm.tmc.edu:8088/quality/info/genbank_annotation.html.

QUALSTAT-REPORT-----

----- Summary Statistics -----
Contig length: 128809
Phrap values in estimate: 128476
Average error rate (BCM-Phrap estimate): 3.45519e-05
Fraction of Phrap values less than 40 : 0.00621906
Number of consensus changing edits: 3
Number of N's in consensus: 0

----- Consensus changing edits -----

Position Original Context Edited Context
93661 gggatcttaa(n)atgggtttgt gggatcttaa(a)atgggtttgt
94679 gctccctta(n)ataaaatgat gctccctta(a)ataaaatgat
119599 aaaaaaaaa(n)gaattaccga aaaaaaaaa(a)gaattaccga

----- Distribution of Quality < 40 Bases -----

# bases	Phrap Value Range									
	5	10	15	20	25	30	35	40		
5001										
4501										
4001										
3501										
3001										
2501										
2001										
1501										
1001										
501										
01		*	*	*	*	*	*	*		

Version: 1.01 gxf.
FEATURES
Location/Qualifiers
Source

1. 128809
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/db_xref="taxon:9606"
/chromosome="12"
/clone="RP11-255114"
complement(1..114)
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complement(1..2005)
/note="Overlaps bases 2005..1 of clone AC025257"
/function="Overlaps with adjacent clone AC025257"
256..327
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264..1339
/rpt_family="MER52A"
1850..1903
/rpt_family="(TAA)n"
complement(2108..2229)
/rpt_family="L1MEC"
2258..2281
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3464..3505
/rpt_family="AT_rich"
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complement(4428..4640)
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5950..6723
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6719..11958
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9223..9436
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10417..10594
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/db_xref="dbSTS:7821"
11994..12330
/rpt_family="THER1B"
12920..12976
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complement(13797..14024)
/rpt_family="L1M4"
complement(14409..14518)
/rpt_family="MIR"

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repeat_region      15194..15627
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repeat_region      15971..15996
                    /rpt_family="AT_Rich"
repeat_region      16346..16396
                    /rpt_family="(TTTTA)n"
repeat_region      complement(16397..16704)
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repeat_region      17398..17434
                    /rpt_family="AT_Rich"

Query Match          42.0%; Score 37.8; DB 9; Length 128809;
Best Local Similarity 64.0%; Pred. No. 29;
Matches 57; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Oy   1 atgaataaatattctcaagttagtgcgttaattttatgcttcattgttcattgtt 60
DB 65107 ATGGAATTAAATGTGCACAGCTACTCTTAATTTAAATAATCCAGATTCATTATAAT 65048
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |
Oy   61 ggaagcataagtcacatgaagaacagcagtc 89
DB 65047 GGAAGAATGAGAAAATAAGAAAAATGTC 65019
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 7
AC079739     152816 bp      DNA             HTG           29-SEP-2000
LOCUS       Homo sapiens chromosome YUK clone CTD-2012B22, WORKING DRAFT
DEFINITION  AC079739
VERSION     AC079739.2 GI:10305265
KEYWORDS    HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE      human.
ORGANISM    Homo sapiens
            Eumetazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 152816)
AUTHORS    Waterston,R.H.
TITLE       The sequence of Homo sapiens clone
JOURNAL    Unpublished
REFERENCE   2 (bases 1 to 152816)
AUTHORS    Waterston,R.H.
TITLE       Direct Submission
JOURNAL    Submitted (10-SEP-2000) Genome Sequencing Center, Washington
            University School of Medicine, 4444 Forest Park Parkway, St. Louis,
            MO 63108, USA
COMMENT     Mo Sep 26, 2000 this sequence version replaced gi:10047917.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site:http://genome.wustl.edu/gsc/index.shtml
Project Information -----
Center project name:H_MS2012B22
Summary Statistics -----
Sequencing vector: MJ3, 95%
Sequencing vector: plasmid, 5%
Chemistry: Dye-primer ET, 95% of reads
Chemistry: Dye-terminator Big Dye, 5% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 130608 bases at least Q40
Consensus quality: 137552 bases at least Q30
Consensus quality: 141453 bases at least Q20
Insert size: 150000; agarose-fp
Quality coverage: 3.13 in Q20 bases; agarose-fp
Quality coverage: 3.30 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 34 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
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		* This record will be updated with the finished sequence	
		* as soon as it is available and the accession number will	
		* be preserved.	
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*	1156	1255:	gap of unknown length
*	1256	2579:	contig of 1324 bp in length
*	2580	2679:	gap of unknown length
*	2680	3782:	contig of 1103 bp in length
*	3783	3882:	gap of unknown length
*	3883	5263:	contig of 1381 bp in length
*	5264	5363:	gap of unknown length
*	5364	7272:	contig of 1909 bp in length
*	7273	7372:	gap of unknown length
*	7373	8845:	contig of 1473 bp in length
*	8846	8945:	gap of unknown length
*	8946	10943:	contig of 1998 bp in length
*	10944	11043:	gap of unknown length
*	11044	12586:	contig of 1543 bp in length
*	12587	12686:	gap of unknown length
*	12687	14738:	contig of 2052 bp in length
*	14739	14838:	gap of unknown length
*	14839	16585:	contig of 1747 bp in length
*	16586	16685:	gap of unknown length
*	16686	18458:	contig of 1773 bp in length
*	18459	18558:	gap of unknown length
*	18559	20559:	contig of 2001 bp in length
*	20560	20559:	gap of unknown length
*	20660	23709:	contig of 3050 bp in length
*	23710	23809:	gap of unknown length
*	23810	25811:	contig of 2002 bp in length
*	25812	25911:	gap of unknown length
*	25912	28324:	contig of 2413 bp in length
*	28325	28424:	gap of unknown length
*	28425	33027:	contig of 4603 bp in length
*	33028	33127:	gap of unknown length
*	33128	37234:	contig of 4107 bp in length
*	37235	37334:	gap of unknown length
*	37335	41889:	contig of 4555 bp in length
*	41890	41989:	gap of unknown length
*	41990	45247:	contig of 3238 bp in length
*	45248	45347:	gap of unknown length
*	45348	50669:	contig of 5332 bp in length
*	50670	50769:	gap of unknown length
*	50770	54317:	contig of 3548 bp in length
*	54318	54417:	gap of unknown length
*	54418	58856:	contig of 4439 bp in length
*	58857	58956:	gap of unknown length
*	58957	63667:	contig of 4711 bp in length
*	63668	63767:	gap of unknown length
*	63768	68619:	contig of 4852 bp in length
*	68620	68719:	gap of unknown length
*	68720	73963:	contig of 5244 bp in length
*	73964	74063:	gap of unknown length
*	74064	78512:	contig of 4449 bp in length
*	78513	78612:	gap of unknown length
*	78613	85002:	contig of 6390 bp in length
*	85003	85102:	gap of unknown length
*	85103	90498:	contig of 5396 bp in length
*	90499	90598:	gap of unknown length
*	90599	98523:	contig of 7925 bp in length
*	98524	98624:	gap of unknown length
*	98624	106597:	contig of 7974 bp in length
*	106598	106697:	gap of unknown length
*	106698	114088:	contig of 7351 bp in length
*	114089	114188:	gap of unknown length
*	114189	123937:	contig of 9749 bp in length
*	123938	124037:	gap of unknown length
*	124038	133862:	contig of 9825 bp in length
*	133863	133962:	gap of unknown length
*	133963	152816:	contig of 18654 bp in length.
		Location/Qualifiers	
		1..152816	
FEATURES		/organism="Homo sapiens"	
source			

/organism="Homo sapiens"

```

Query Match Similarity   40.4% ; Score 36.4 ; DB 2; Length 152816;
Best Local Similarity    64.0%; Pred.No.57;
Matches      55; Conservative     0; Mismatches     31; Indels       0; Gaps        0;

Qy          3 gaataaatattcctaaagttactgtaatcttcttcgaacgtttccattcgtagtg 62
            | ||||| ||||| ||||| || ||||| ||||| ||||| |||||
Db  42020 gaaaatatatttatTTAAATATTACGCTTTCrNATTTTTTAAGTTTTTGCGITTTTC 42079

Qy         63 agcaataagtccaatgaaagaacaagt 88
            + | || | | | | | | | | | |
Db  42080 TACCACAAGAATCCTATAAGCAATGG 42105

RESULT# 8
LOCUS AC022199/C
DEFINITION Homo sapiens chromosome 17 clone RP11-145G19 map 17, WORKING DRAFT SEQUENCE, 12 unordered pieces.
ACCESSION AC022199
VERSION AC022199.2 GI:7417783
KEYWORDS HTG; HTGS_PHASEI; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE Birren,B., Linton,L., Nusbaum,C., Landier,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barna,N., Beckery,R., Bedalov,A., Boguslavsky,I., Bouckgalter,B., Brown,A., Burkett,G., Castle,A., Choepel,Y., Colangelo,M., Collins,S., Collamore,A., Cooke,P., DeRellano,K., Dewar,K., Domino,M., Doyle,M., Festeron,J., Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos-B., Heaford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., Landers,T., Lehoczyk,J., Levine,R., Lieu,C., Liu.G., Locke.K., Macdonald,P., Marquis.N., McEwan.P., McGurk.A., McKernan,K., McPheters.R., Meldrim.J., Menius.L., Morrow.J., Naylor,J., Norman,C.H., O'Connor.T., O'Donnell.P., Olivari.I.M., Peterson,K., Pieree,N., Pisani,C., Pollara,V., Raymond.C., Riley.R., Rothman.D., Roy,A., Santos.R., Severy,P., Spencer.B., Stange-Thomann,N., Stojanovic-N., Subramanian,A., Talmas,J.J., Tesfaye.S., Theodore,J., Tirrell,A., Vassiliou.H., Viel,R., Vo.A., Wu.X., Wyman,D., Ye.W.J., Zimmer,A. and zody.M.
DIRECT SUBMISSION Submitted (26-JAN-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Apr 5, 2000 This sequence version replaced by gi:6759152.
All repeats were identified using RepeatMasker:
Smit, A.F.A & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center -----
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information -----
Center project name: I5174
Center clone name: I45-G_19

----- Summary Statistics -----
Sequencing vector: MI3, M7815, 100% of reads
Assembly: Dye-terminator Big Dye; 100% of reads
Assembler program: Phrap; version 0.960731
Consensus quality: 157408 bases at least Q40
Consensus quality: 161068 bases at least Q30
Consensus quality: 162457 bases at least Q20
Insert size: 165000; agarose-gel
Insert size: 164076; sum-of-contigs
```



```
misc_feature      /organism="Homo sapiens"
                  /db_xref="taxon:9606"
                  /clone="RP11-11E12"
                  /clone_11b="RP11 Human Male BAC"
                  1..4427
misc_feature      /note="assembly-fragment
                  clone_end:SP6
                  vector_side:left"
                  4428..6681
misc_feature      /note="assembly-fragment"
                  6982..9727
misc_feature      /note="assembly-fragment"
                  9828..12474
misc_feature      /note="assembly-fragment"
                  12575..14864
misc_feature      /note="assembly-fragment"
                  14965..18876
misc_feature      /note="assembly-fragment"
                  18977..60052
misc_feature      /note="assembly-fragment"
                  60153..69381
misc_feature      /note="assembly-fragment"
                  69482..77833
misc_feature      /note="assembly-fragment"
                  77934..86115
misc_feature      /note="assembly-fragment"
                  86216..98453
misc_feature      /note="assembly-fragment"
                  98554..116358
misc_feature      /note="assembly-fragment"
                  116459..134093
misc_feature      /note="assembly-fragment"
                  134194..162917
misc_feature      /note="assembly-fragment"
                  163018..167322
```

1	43166	contig of	43166	bp in length
43367	63703	contig of	19437	bp in length
62804	84476	contig of	21673	bp in length
84577	106668	contig of	32092	bp in length
106769	128319	contig of	19551	bp in length
126620	141481	contig of	15062	bp in length
141582	153916	contig of	12335	bp in length

154017 163781 contig of 9765 bp in length
 163882 170760 contig of 6879 bp in length
 170861 176864 contig of 6004 bp in length
 176965 183161 contig of 6197 bp in length
 183262 185847 contig of 2586 bp in length
 185948 189771 contig of 3824 bp in length.
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 13 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 43166: contig of 43166 bp in length
 * 43167 43266: gap of 100 bp
 * 43267 62703: contig of 19437 bp in length
 * 62704 62803: gap of 100 bp
 * 62804 84476: contig of 21673 bp in length
 * 84477 84576: gap of 100 bp
 * 84577 106668: contig of 22092 bp in length
 * 106669 106768: gap of 100 bp
 * 106769 126319: contig of 19551 bp in length
 * 126320 126419: gap of 100 bp
 * 126420 141481: contig of 15062 bp in length
 * 141482 141581: gap of 100 bp
 * 141582 153916: contig of 12335 bp in length
 * 153917 154016: gap of 100 bp
 * 154017 163781: contig of 9765 bp in length
 * 163782 163881: gap of 100 bp
 * 163882 170760: contig of 6879 bp in length
 * 170761 170860: gap of 100 bp
 * 170861 176864: contig of 6004 bp in length
 * 176865 176964: gap of 100 bp
 * 176965 183161: contig of 6197 bp in length
 * 183162 183261: gap of 100 bp
 * 183262 185847: contig of 2586 bp in length
 * 185848 185947: gap of 100 bp
 * 185948 189771: contig of 3824 bp in length.

FEATURES

source
 1. 189771
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="11"
 /map="11g"
 /clone="RP11-480C22"
 1. 43166
 /note="assembly-fragment"
 43267. 62703
 /note="assembly-fragment"
 62804. 84476
 /note="assembly-fragment clone_end:17 vector_side:left"
 84577. 106668
 /note="assembly-fragment"
 106769. 126319
 /note="assembly-fragment"
 126420. 141481
 /note="assembly-fragment"
 141582. 153916
 /note="assembly-fragment"
 154017. 163781
 /note="assembly-fragment"
 163882. 170760
 /note="assembly-fragment"
 170861. 176864
 /note="assembly-fragment clone_end:SP6 vector_side:left"
 176965. 183161
 /note="assembly-fragment"
 183262. 185847
 /note="assembly-fragment"
 185948. 189771
 /note="assembly-fragment"
 misc_feature
 56729 a 37254 c 36581 g 58007 t 1200 others

ORIGIN

Query Match 40.4%; Score 36.4; DB 2; Length 189771;
 Best Local Similarity 64.0%; Pred. No. 57;
 Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

OY 3 gaataaatttttaagttacgtatattatglttcacgtttcttatgttg 62
 Db 140789 GAAAAATATTATTAATAATATTACGTCTTCTATATATTATTTCTGCTTTTC 140730
 OY 63 agcaataagtcacatgaaagcaagtg 88
 Db 140729 TACCAACAGCAATCTTATAGTCAATG 140704

RESULT 11
 AC074180/c DNA HTG 17-AUG-2000
 LOCUS
 DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 193415)
 Waterston,R.H.
 The sequence of Homo sapiens clone
 Unpublished
 2 (bases 1 to 193415)
 Waterston,R.H.
 Direct Submission
 Submitted (15-JUL-2000) Genome Sequencing Center, Washington
 University School of Medicine, 4444 Forest Park Parkway, St. Louis,
 MO 63108, USA
 On Aug 17, 2000 this sequence version replaced gi.9211440.

COMMENT

----- Genome Center -----
 Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: http://genome.wustl.edu/gsc/index.shtml
 ----- Project Information -----
 Center project name: H NH0480C22
 ----- Summary Statistics -----
 Sequencing vector: M13; 2x
 Sequencing vector: plasmid; 0x
 Chemistry: Dye-primer ET; 2x of reads
 Chemistry: Dye-terminator Big Dye; 0x of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 187074 bases at least Q40
 Consensus quality: 192835 bases at least Q30
 Consensus quality: 195619 bases at least Q20
 Insert size: 182000; agarose-1p
 Insert size: 195112; sum-of-contigs
 Quality coverage: 7.73 in Q20 bases; agarose-1p
 Quality coverage: 7.34 in Q20 bases; sum-of-contigs
 ----- NOTE: This is a 'working draft' sequence. It currently
 * consists of 10 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
 1 3180: contig of 3180 bp in length
 * 3181 3280: gap of unknown length
 * 3281 12431: contig of 9151 bp in length
 * 12432 12531: gap of unknown length
 * 12532 23140: contig of 10609 bp in length


```

* 23141 23240: gap of unknown length
* 23241 34935: contig of 11695 bp in length
* 34936 35035: gap of unknown length
* 35036 50440: contig of 15405 bp in length
* 50441 50540: gap of unknown length
* 50541 71299: contig of 20759 bp in length
* 71300 71399: gap of unknown length
* 71400 96045: contig of 24646 bp in length
* 96046 96145: gap of unknown length
* 96146 124103: contig of 27958 bp in length
* 124104 124203: gap of unknown length
* 124204 160061: contig of 35858 bp in length
* 160062 160161: gap of unknown length
* 160162 193415: contig of 33254 bp in length.

```

```

FEATURES
source
1. 193415
Location/Qualifiers

```

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="11"
/clone="RP11-480C22"
1. 3180
misc_feature
/feature="assembly_name:Contig21"
3281..12431

```

```

misc_feature
/feature="assembly_name:Contig22"
12532..23140
misc_feature
/feature="assembly_name:Contig23"
23241..34935

```

```

misc_feature
/feature="assembly_name:Contig24"
35036..50440
misc_feature
/feature="assembly_name:Contig25"
50541..71299

```

```

misc_feature
/feature="assembly_name:Contig26"
71400..96045
misc_feature
/feature="assembly_name:Contig27"
96146..124103

```

```

misc_feature
/feature="assembly_name:Contig28"
124204..160061
misc_feature
/feature="assembly_name:Contig29"
160162..193415

```

```

misc_feature
/feature="assembly_name:Contig30"
clone_end:SP6
vector_side:right"

```

```

BASE COUNT 57779 a 38131 c 37181 g 59416 t 908 others
ORIGIN

```

```

Query Match 40.4%; Score 36.4; DB 2; Length 193415;
Best Local Similarity 64.0%; Pred. No. 57;
Matches 55; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

```

```

QY 3 gaaataattattcaagttactgtatatttatgtttcattcttctatgttg 62
DB 21903 GAAATAATTATTCAAGTTACTGTATATTATGTTTATGTTTCTGTTTTC 21844
QY 63 agcataagtcataagaaagcaagt 88
DB 21843 TACCAAGATCTTATAGTCAATG 21818

```

```

RESULT 12
LOCUS CENS01BGA 148 bp mRNA PLN 02-SEP-1999
DEFINITION Botrytis cinerea strain T4 cDNA library under conditions of
nitrogen deprivation.
ACCESSION AL114194
VERSION AL114194.1 GI:5828813
KEYWORDS cDNA library; nitrogen deprivation.
SOURCE Botryotinia fuckeliana
ORGANISM Eukaryota; Fungi; Ascomycota; Pezizomycotina; Leotiomycetes;
Helotiales; Sclerotiniaceae; Botryotinia.
REFERENCE 1 (bases 1 to 148)
Bilton,F., levis,C., Fortini,D., Pradler,J.M. and Brygoo,Y.

```

```

TITLE Direct Submission
JOURNAL Submitted (01-SEP-1999) Phytopathologie, INRA, route de St Cyr,
78026 Versailles, France
REFERENCE 2 (bases 1 to 148)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (01-SEP-1999) Genoscope - Centre National de Sequencage :
CP 5706 91057 Evry cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)

```

COMMENT

The cDNA library to be analyzed within the framework of this project was created using a botrytis cinerea strain which was grown under conditions of nitrogen deprivation, which is the normal situation for B. cinerea during its development on its host plant. The library was produced in an oriented direction, in the pBSII vector.

```

FEATURES
source
1. 148
/organism="Botryotinia fuckeliana"
/strain="T4"
/db_xref="taxon:40559"
/feature="Genoscope sequence ID : W52A031"

```

```

BASE COUNT 33 a 17 c 25 g 73 t
ORIGIN

```

```

Query Match 39.38; Score 35.4; DB 8; Length 148;
Best Local Similarity 66.2%; Pred. No. 1e+02;
Matches 51; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

```

```

QY 11 ttattcaagttactgtatatttatgtttcattcttctatgttgagcaataa 70
DB 45 TTATTATTTTTCATATATTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTT 104
QY 71 gtccaatgaagcaagt 87
DB 105 GAGCATCAACATTAAGT 121

```

```

RESULT 13
AF099924/c 33460 bp DNA INV 09-AUG-2001
LOCUS Caenorhabditis elegans cosmid K07A9, complete sequence.
DEFINITION AF099924
ACCESSION AF099924
VERSION AF099924.2 GI:6671811
KEYWORDS HTG.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
Rhabditioidea; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 33460)

```

```

REFERENCE 1 The C. elegans Sequencing Consortium.
Genome sequence of the nematode C. elegans: a platform for
investigating biology. The C. elegans Sequencing Consortium
Science. 282 (5396), 2012-2018 (1998)
JOURNAL 99069613
MEDLINE 2 (bases 1 to 33460)
REFERENCE Davidson,S. and O'Neal,D.
AUTHORS The sequence of C. elegans cosmid K07A9
unpublished
JOURNAL 3 (bases 1 to 33460)
REFERENCE Waterston,R.
AUTHORS Direct Submission
JOURNAL 4 (bases 1 to 33460)
REFERENCE Waterston,R.
AUTHORS Direct Submission
JOURNAL 5 (bases 1 to 33460)
REFERENCE Waterston,R.
AUTHORS Direct Submission
JOURNAL Submitted (27-MAY-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

```

REFERENCE 6 (bases 1 to 33460)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (05-JAN-2000) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
7 (bases 1 to 33460)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (09-AUG-2001) Department of Genetics, Washington University, Genome Sequencing Center, 4444 Forest Park Avenue, St. Louis, MO 63110, USA
On Jan 5, 2000 this sequence version replaced gi:3790764.
Submitted by:
Genome Sequencing Center
Department of Genetics, Washington University
St. Louis, MO 63110, USA, and
Sanger Centre, Hinxton Hall
Cambridge CB10 1RO, England
email: rwenematode.wustl.edu and joes@sanger.ac.uk

COMMENT
On Jan 5, 2000 this sequence version replaced gi:3790764.
Submitted by:

NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighboring submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one m13 subclone.

NEIGHBORING COSMID INFORMATION

The 5' clone is Y38C1B, 200 bp overlap; 3' clone is W03A6, 200 bp overlap. Actual start of this clone is at base position 1 of CELK07A9; actual end is at 11466 of CELM03A6.

NOTES:

Coding sequences below are predicted from computer analysis, using the program Genefinder(P. Green and L. Hillier, ms in preparation). Location/Qualifiers

FEATURES
source

1..33460
/organism="Caenorhabditis elegans"

/strain="Bristol N2"

/db_xref="taxon:6239"

/chromosome="IV"

/clone="K07A9"

96..9167

/gene="K07A9.2"

/note="cmk-1"

join(96..200,6063..6181,7483..7639,7732..7997,8434..8537,8587..8756,9042..9167)

/gene="K07A9.2"

/note="C. elegans Ca2+/calmodulin-dependent protein kinase I (GB:AB021864); contains similarity to Pfam domain PF00069 (pkinese), Score=262.3, E-value=2e-75; N=1; coded for by the following C. elegans cDNAs: YK306d5.3, YK306d5.5, AB021864"

/codon_start=1

/product="Hypothetical protein K07A9.2"

/protein_id="AAZ23187.1"

/db_xref="GI:6671812"

/translation="MPLFKRRDGGSPAPNATIREKYDFRDVLGTGAFSGVFLAEKSDACQMYAVKCIDKALKGKESLENEIKVRLKLNHN1VOLFEDYDKQVYLMELVLTGGELPRIVAKSGSYTEODASNLIRQVLEAVGEHMDKGVNHRDLKPNLLIYNDDESKIMTSDRGISTKSTEDSGMATACGTPGVAVPEVLOOKYGVKAVDVSGIAYILICGYPPTDESDANIFAQIIEGVEFPAPYWDQISDSAKDPTIHLMDCCDPARFECODALSHPWISGNATYTHDIGIYAVHLKSLARKNNKATNAALAIROLMLRLSSNSNLTQKQASQOQPEPPPAFPA"

/gene="K07A9.3"

10199..11708

/gene="K07A9.3"

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

gene

CDS
join(10199..10231,10572..10658,10709..10810,11291..11470,11541..11708)
/gene="K07A9.3"

/codon_start=1
/product="Hypothetical protein K07A9.3"

/protein_id="AAC68811.1"

/db_xref="GI:3790766"

/translation="MDASTSEFATDFGIISNEKFSGLDPFGRTDFLRIRYDKAKNKSIRKSTHVDNFRAYITIOVLHGKIVVYDRKLYVNNYLGFLINQPMVFAQLSEISIOKRRKKTKVSPSPNFRASLIPROKIGIONSQFTCEFSIPIFSCGSLTEGVLVFTFNIAPIQGNINRDLMLVLPDEASLS"

BASE COUNT 9970 a 6275 c 6181 g 11034 t
ORIGIN

Query Match 39.3%; Score 35.4; DB 3; Length 33460;
Best Local Similarity 63.5%; Pred. No. 96;
Matches 54; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

QY 66 aataagctcaagtctactgtaatttctatgcttcatgcttcttcttcttctgagc 65
DB 27539 AAAAGCTTTTGAATAATTTCGAAATTTTAATTCACATTTCTTCTTTTAAAG 27480

RESULT 14
AF067612/c
LOCUS AF067612 Caenorhabditis elegans cosmid F55A8, complete sequence.
DEFINITION AF067612
ACCESSION AF067612
VERSION AF067612.2 GI:4982487
KEYWORDS HTG.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabdilitida; Rhabdilitidae; Rhabdilitidae; Pelodierinae; Caenorhabditis.

REFERENCE 1 (bases 1 to 69431)
AUTHORS The C. elegans Sequencing Consortium.
TITLE Genome sequence of the nematode C. elegans: a platform for investigating biology. The C. elegans Sequencing Consortium
JOURNAL Science 282 (5396), 2012-2018 (1998)
MEDLINE 99069613
REFERENCE 2 (bases 1 to 69431)
AUTHORS Langston, Y., Wohldmann, P. and Duckels, G.
TITLE The sequence of C. elegans cosmid F55A8
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 69431)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 4 (bases 1 to 69431)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (22-MAY-1998) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
REFERENCE 5 (bases 1 to 69431)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (03-JUN-1999) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
REFERENCE 6 (bases 1 to 69431)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (03-JUL-2001) Department of Genetics, Washington University, Genome Sequencing Center, 4444 Forest Park Avenue, St. Louis, MO 63110, USA
On Jun 3, 1999 this sequence version replaced gi:3193146.
Submitted by:
Genome Sequencing Center
Department of Genetics, Washington University

St. Louis, MO 63110, USA, and
Sanger Centre, Hinxton Hall
Cambridge CB10 1RQ, England
email: rwenemate@wustl.edu and jesus@sanger.ac.uk

NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections or longer because we provide a small overlap between neighboring submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one ml3 subclone.

NEIGHBORING COSMID INFORMATION

The 5' clone is W03A6, 22600 bp overlap; 3' clone is Y48A5B, 2100 bp overlap. Actual start of this clone is at base position 22401 of CELF55A8; actual end is at 67531 of CELF55A8.

NOTES:

Coding sequences below are predicted from computer analysis, using the program GeneFinder (P. Green and L. Hillier, ms in preparation). Location/Qualifiers

FEATURES

source

gene

CDS

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/db_xref="taxon:6239"
/chromosome="IV"
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29221. 29854,31878. 31973)
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(pkinaase_C, Score=16.7, E-value=0.00024, N=1); coded for
by the following C. elegans cDNAs: YK2f8.5, YK4g10.3,
YK17d12.3, YK30f8.5, YK41a5.3, YK52d1.3, YK52d1.5,
YK55a6.3, YK55a6.5, YK55f2.3, YK55f2.5, YK81h11.3,
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YK325g6.5, YK328c10.5, YK341c9.5, YK349e8.5, YK385d7.5,
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gene

CDS

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(pkinaase_C, Score=16.7, E-value=0.00024, N=1); coded for by
the following C. elegans cDNAs: YK2f8.5, YK4g10.3,
YK4g10.5, YK812.5, YK17d12.3, YK30f8.5, YK41a5.3,
YK41a5.5, YK52d1.3, YK52d1.5, YK55a6.3, YK55a6.5,
YK55f2.3, YK55f2.5, YK81h11.3, YK81h11.5, YK87g1.3,
YK87g1.5, YK87g12.3, YK87g12.5, YK193d8.5, YK243f5.5,
YK285h5.5, YK317g2.5, YK348d12.5, YK349e8.5, YK385d7.5,
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BASE COUNT 21037 a 13563 c 12501 g 22330 t
ORIGIN

```

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Query Match 39.3%; Score 35.4; DB 3; Length 69431;
Best Local Similarity 63.5%; Pred. No. 95;
Matches 54; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

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||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 4004 AAAAGCTTTTGAATTTTTCAGAAATTTTAAATTTCCACATTTTCATTTTAAAGC 3945

QY 66 aataagttcaatgaagaagtga 90
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 3944 CAGAAATTTCTGGGATGCCAGTACA 3920

RESULT 15
AC006364/c 132319 bp DNA PRI 21-DEC-1999
DEFINITION Homo sapiens BAC clone GSI-146J4 from 7q31.1-31.3, complete
sequence.
AC006364
VERSION AC006364.3 GI:4753253
KEYWORDS HTG.

```

SOURCE ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE TITLE 1 (bases 1 to 132319)
Sulston, J.E. and Waterston, R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)

REFERENCE MEDLINE 99063792
2 (bases 1 to 132319)
Ryan, E., Bauer, C., Tuccil, S. and Spalding, L.
The sequence of Homo sapiens BAC clone GSI-146J4

REFERENCE JOURNAL Unpublished
3 (bases 1 to 132319)
Waterston, R.H.
Direct Submission
Submitted (11-JAN-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

REFERENCE JOURNAL 4 (bases 1 to 132319)
Waterston, R.H.
Direct Submission
Submitted (05-MAY-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

REFERENCE JOURNAL 5 (bases 1 to 132319)
Waterston, R.
Direct Submission
Submitted (11-SEP-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On May 5, 1999 this sequence version replaced g1:4337266.

REFERENCE JOURNAL 6 (bases 1 to 132319)
Waterston, R.
Direct Submission
Submitted (21-DEC-1999) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On May 5, 1999 this sequence version replaced g1:4337266.

COMMENT

Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@wustl.wustl.edu

Summary Statistics

Center project name: H_GSI46J04

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
The sequence of this clone was established as part of a mapping and sequencing collaboration between the NHGRI Chromosome 7 Mapping Project (Eric D. Green, Director), John D. McPherson in the Department of Genetics (Washington University), and the Washington University Genome Sequencing Center. For additional information about the map position of this sequence, see
http://www.nhgri.nih.gov/DIR/CTB/CHR7, send
mailto:egreen@nhgri.nih.gov, or see http://genome.wustl.edu/gsc

SOURCE INFORMATION:
This clone is from the first BAC library from Genome Systems, Inc.
(http://www.genomesystems.com).
Cell line: lymphoblastoid
Haplotypes: two

VECTOR: pBelBAC
Selection: chloramphenicol
NEIGHBORING SEQUENCE INFORMATION:
The actual start of this clone is at base position 1 of GSI-146J4.
Actual end is at 132319 of GSI-146J4.
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2512. 2783
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EGCIEPKOD"
11766. 12114

misc_feature

Search completed: January 24, 2002, 02:27:00
Job time: 4473 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 01:51:28 ; Search time 2272.52 Seconds
(without alignments)
425.572 Million cell updates/sec

Title: US-09-531-438-4
Perfect score: 90
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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 11351937 seqs, 5372889281 residues

Total number of hits satisfying chosen parameters: 22703874

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

EST:*
1: em_estfun:*
2: em_esthum:*
3: em_estlin:*
4: em_estlom:*
5: em_estlpl:*
6: em_estda:*
7: em_estro:*
8: em_estov:*
9: em_hlc:*
10: gb_estl:*
11: gb_est2:*
12: gb_hlc:*
13: gb_gss:*
14: em_gss_fun:*
15: em_gss_hum:*
16: em_gss_inv:*
17: em_gss_pln:*
18: em_gss_pro:*
19: em_gss_rtd:*
20: em_gss_vrt:*
21: em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	37.8	42.0	561	13	AO533830 RPCI-11-3
C 2	35.6	39.6	302	10	AV247937
C 3	34.8	38.7	447	13	B67487 T25M7TR TAM
C 4	34.8	38.7	482	13	AO968516 LERJDO3TR
C 5	34.8	38.7	488	13	AO968397 LERJC21TR
C 6	34.8	38.7	497	13	AO958266 LERAW47TR
C 7	34.8	38.7	498	13	B77731 T28M11TR TA
C 8	34.8	38.7	623	13	B30114 T26F22TRB T
C 9	34.8	38.7	627	13	B78178 T31J11TR TA
C 10	34.8	38.7	640	13	AO958844 LEREF16TR
C 11	34.8	38.7	643	13	B98363 T24C11TRB T
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C 14	34.8	38.7	743	13	AO958265 LERAW47TR
C 15	34.8	38.7	798	13	AO956484 LERAK26TR
C 16	34.4	38.2	607	13	AZ522516 207PDC02
C 17	34.4	38.2	638	13	AZ523727 222PBH02
C 18	34.2	38.0	923	13	BH132631 ENTOP7OTR
C 19	34.2	37.8	871	13	AZ680048 ENTRK43TR
C 20	33.8	37.6	867	13	CNS00CX5 AL060052 DROSOPH11
C 21	33.6	37.3	240	10	AU072517 AU072517
C 22	33.6	37.3	240	10	AU072714 AU072714
C 23	33.6	37.3	306	11	C24362 C24362 Dict
C 24	33.6	37.3	413	10	AV760889 AV760889
C 25	33.6	37.3	376	13	AO798320 HS_3160_A
C 26	33.6	37.3	552	13	AZ065684 EST502514
C 27	33.6	37.3	676	13	AO635998 RPCI-11-4
C 28	33.6	37.3	913	13	BH132369 ENTNH76TR
C 29	33.4	37.1	432	10	AA149033 ZO3B12.S
C 30	33.2	36.9	463	11	BG603509 EST502599
C 31	33.2	36.9	526	11	BG603424 EST502514
C 32	33.2	36.9	652	11	BG603423 EST502513
C 33	33.2	36.7	1101	13	CNS00K05 AL076967 DROSOPH11
C 34	32.8	36.4	420	11	Z38090 Z38090 ATT54230 Ve
C 35	32.8	36.4	473	13	AO001587 CIT-HSP-2
C 36	32.8	36.4	589	13	AO026620 CIT-HSP-2
C 37	32.8	36.4	843	13	AZ549410 ENTF80TF
C 38	32.8	36.4	899	13	BH136018 ENTOL96TF
C 39	32.8	36.4	944	13	AZ543633 ENTDL7TR
C 40	32.8	36.4	949	13	AL109198 DROSOPH11
C 41	32.8	36.4	958	13	AZ687864 ENTH82TR
C 42	32.6	36.2	510	13	AZ384026 IM0141G19
C 43	32.6	36.2	553	13	AO698845 HS_5557_B
C 44	32.4	36.0	366	13	BH014409 TOSBIA2TR
C 45	32.4	36.0	589	13	AZ247746 RPCI-23-9

ALIGNMENTS

RESULT 1	AO533830/c	561 bp	DNA	GSS	18-MAY-1999
LOCUS	RPCI-11-379L24.TV	RPCI-11	Homo sapiens genomic clone RPCI-11-379L24		
DEFINITION	, DNA sequence.				
ACCESSION	AO533830				
VERSION	AO533830.1	GI:4845520			
KEYWORDS	GSS.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter,J.C.				
TITLE	Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready				
JOURNAL	Map Building				
COMMENT	Unpublished (1997)				
	Other-GSSs: RPCI-11-379L24.TJ				
	Contact: Shaying Zhao, William Nierman, Mark Adams				
	Department of Eukaryotic Genomics				
	The Institute for Genomic Research				
	9712 Medical Center Dr., Rockville, MD 20850				
	Tel: 301 838 0200				
	Fax: 301 838 0208				
	Email: hbe@tigr.org				
	Clones are derived from the human BAC library RPCI-11. For BAC				
	library availability, please contact Pieter de Jong				
	(pieter@jeong.med.buffalo.edu). Clones may be purchased from				
	BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from				
	Research Genet cs (info@resgen.com). BAC end search page:				
	http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.				
	Seq primer: T7				
	Class: BAC ends.				
	Location/Qualifiers				

FEATURES

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/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC111 Human Male BAC Library"

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Best Local Similarity 73.8%; Fred. No. 2.6e+02;
Matches 48; Conservative 0; Mismatches 17; Indels 0; Gaps 0

Oy      4      aaaaattattccaaattctactgaatttaattgaatttcacgtttcttattgttga 63
||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      428      AAGAGATATTATTCATGTTCTTGCTCTTTTTCATTTCTTTTATTTGATGA 369

Oy      64      gcaat 68
|||
Db      368      GAAAT 364

RESULT 2
AV247937      302 bp      mRNA      EST      04-NOV-1999
AV247937      RIKEN full-length enriched, 0 day neonate head Mus
musculus cDNA clone 4833401E05 3' similar to X64411 R.norvegicus
RNA for 100 kDa protein, mRNA sequence.
AV247937
AV247937.1 GI:6235396
EST.
house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 302)
Kono,H., Aizawa,K., Akahira,S., Akiyama,J., Carninci,P., Endo,T.,
Fukunoh,S., Fukunishi,Y., Hara,A., Hayatsu,N., Hirazane,T., Hori,F.,
Ishii,Y., Ishikawa,T., Itoh,M., Izawa,M., Kadota,K., Kagawa,I., Ka-
,C., Kawai,T., Kikuchi,N., Kojima,Y., Koya,S., Kusakabe,M.,
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Owa,C., Ozawa,Y., Saito,H., Sano,M., Sato,K., Shibata,K., Shibata
,Y., Shigemoto,Y., Shiraki,T., Sogabe,Y., Sugahara,Y., Suzuki,H.,
Suzuki,H., Takahashi,F., Tateo,M., Tominaga,N., Tsunoda,Y.,
Watanishi,A., Watanabe,S., Yamamura,T., Yasunishi,A., Yokota,T.,
Yoshiki,A., Yoshino,M., Muramatsu,M. and Hayashizaki,Y.
RIKEN Mouse ESTs (Kono,H., et al. 1999)
Unpublished (1999)
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Shuho-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Sasaki,N., Izawa,M., Watanishi,M., Ozawa,K., Tanaka,T., Yoneda,Y.,
Matsura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and Hayashizaki
,Y.
Transcriptional sequencing: A method for DNA sequencing using RNA
polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
Itoh,M., Kikuchi,T., Akiyama,Y., Shibata,K., Izawa,M., Kawai,J.,
Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki
,Y. and Hayashizaki,Y.
Automated filtration-based high-throughput plasmid preparation
system. Genome Res. 9 (5), 463-470 (1999)
Carninci,P. and Hayashizaki,Y.

```

High-efficiency full-length cDNA cloning: *Methods Enzymol.* 303:19-44 (1999)
Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES	
SOURCE	Location/Qualifiers 1..302 /organism="Mus musculus" /strain="C57BL/6J" /db_xref="taxon:10090" /clone="483401E05" /clone_id="RIKEN full-length enriched, 0 day neonate head" /sex="mixed" /tissue_type="head" /dev_stage="0 day neonate" /lab_host="DH10B" /note="Site_1: SalI; Site_2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5', GAGAGAGAGAGATCCACGACTCTTTTGTTCCTTTTTCVN 3'], cDNA was prepared by using triethanolamine thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 10.0 and subtraction to Rot = 100.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGAGATTTCGAGTTTAATAATTAATCCCCCCCC 3']. cDNA was cloned into the xhoI and BamHI sites. Vector: a modified pluescript KS(+) after bulk excision from Lambda FIC I."
BASE COUNT	86 a 58 c 62 g 96 t
ORIGIN	
Query Match	39.6%; Score 35.6; DB 10; Length 302;
Best Local Similarity	64.6%; Pred. No. 8.9e+02;
Matches	53; Conservative 0; Mismatches 23; Indels 0; Gaps 0;
OY	4 aaaaataattccaagtcttaactgtlaattttatglttcacgtttcattatgttgga 63 Db 185 AAAAAAATAATCATCGAGTGTAATTAATTTGGTTCGTGCATCTTTGGCTGGGTGA 126 OY 64 gcaataagtlccaatgaagaaca 85
Db	125 CCTAGAAGTCGGGAGTAAGCAA 104
RESULT 3	
B67487/c	B67487 447 bp DNA GSS 09-DEC-1997
DEFINITION	T25M7R TAMU Arabidopsis thaliana genomic clone T25M7, DNA sequence.
ACCESSION	B67487
VERSION	B67487.1 GI:2666241
KEYWORDS	GSS.
SOURCE	thale cress.
ORGANISM	Arabidopsis thaliana Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE	1 (bases 1 to 447) Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Golden,K., Berry,R., Granger,D., Suh,E., Wilde,C., Adams,M.D. and Venter
AUTHORS	J.D.C.
TITLE	A BAC End Sequence Database for Identifying Minimal Overlaps in Arabidopsis Genomic Sequencing. Update 3
JOURNAL	Unpublished (1997)
COMMENT	Contact: Steve Rounsley Department of Eukaryotic Genomics The Institute for Genomic Research


```

RESULT      6
LOCUS       A0958266
DEFINITION  A0958266 497 bp DNA
ACCESSION   LERAW47.LERA Arabidopsis thaliana genomic clone LERAW47, DNA
VERSION     A0958266
KEYWORDS    A0958266.1 GI:6785967
SOURCE      GSS.
ORGANISM    thale cress.
            Arabidopsis thaliana
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
            1 (bases 1 to 497)
            Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Utterbach,T.,
            Feldblum,T., Liang,F., Creasy,T. and Fraser,C.M.
            Genomic survey sequencing of Landsberg erecta ecotype of
            Arabidopsis thaliana and identification of sequence-based
            polymorphisms
JOURNAL     Unpublished (2000)
COMMENT     Contact: Xiaoying Lin
            The Institute for Genomic Research
            9712 Medical Center Dr., Rockville, MD 20850, USA
            Tel: 301 838 0200
            Fax: 301 838 0208
            Email: atctigr.org
            For additional information, see http://www.tigr.org/tdb/at/at.html
            Similar to A. thaliana chloroplast sequence (GB:AP000423)
            Seq primer: TR
            Class: shotgun.
FEATURES
    source          location/Qualifiers
                     1..497
                     /organism="Arabidopsis thaliana"
                     /strain="Landsberg erecta"
                     /db_xref="taxon:3702"
                     /clone="LERAW47"
                     /clone_lib="LERA"
                     /note="Organ: Leaf; Vector: pHOS1; Total genomic DNA was
                     sheared to 0.9-1 kbp before ligation."
BASE COUNT      144 a                82 c                69 g                202 t
ORIGIN
Query Match      38.7%; Score 34.8; DB 13; Length 497;
Best Local Similarity 65.4%; Pred. No. 9,9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
OY      5. aaaaattttcaaaagttcactgaatttaatttaagtttcacgtttcttattgttgag 64
        ||||| |||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB      337 AAATTAATAATTTGGAAGAGTTTGAATTCCTCTCTGCAATTAATTTCTCTGTTGGGG 396
        ||||| |||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
OY      65 caataagtcacgaagaag 82
        ||||| ||||| |||||
DB      397 TTCTGAAAAAAGAAAG 414
        ||||| ||||| |||||
RESULT      7
B77731/c      498 bp DNA
LOCUS         B77731
DEFINITION    T28N11TF TAMU Arabidopsis thaliana genomic clone T28N11, DNA
ACCESSION     B77731
VERSION       B77731
KEYWORDS      B77731.1 GI:274370
SOURCE        GSS.
ORGANISM      thale cress.
            Arabidopsis thaliana
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
            1 (bases 1 to 498)
            Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Linher,K., Golden,K.
            Berry,K., Granger,D., Sun,E., Wible,C., Adams,M.D. and Venter
            ,J.C.

```

TITLE A BAC End Sequence Database for Identifying Minimal Overlaps in Arabidopsis Genomic Sequencing. Update 3
 JOURNAL Unpublished (1997)
 COMMENT Other_GSSs: T28N117R
 Contact: Steve Rounsley
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: rounsley@tigr.org
 Seq primer: M13-21
 Class: BAC ends
 High quality sequence stop: 498.
 Location/Qualifiers
 1. 498
 /organism="Arabidopsis thaliana"
 /strain="Columbia"
 /db_xref="taxon:3702"
 /clone="T28N11"
 /clone_1lb="TAMU"
 /sex="hermaphrodite"
 /note="Vector: Bel0BAC11; Site_1: HindIII; Site_2: HindIII
 ; Produced by Rod Wing"
 BASE COUNT 210 a 68 c 72 g 148 t
 ORIGIN
 Query Match 38.7%; Score 34.8; DB 13; Length 498;
 Best Local Similarity 65.4%; Pred. No. 9.9e+02;
 Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
 QY 5 aaaaatattcaaaagtctacgtcaatttcatgtttcatgtttctattgttgag 64
 ||| || |||| ||||| | |||| || |||| ||||| ||||| |
 DB 379 AAATATAAAATTGAAAGCTTGATTTCTTCCTCGAATTAATTTTCTCTGTGGGG 320
 ||| || |||| ||||| | |||| || |||| ||||| ||||| |
 QY 65 caataagtcacaatgaag 82
 | | | | |
 DB 319 TTCTGAAAAAAGAAAG 302
 ||| || |||| ||||| | |||| || |||| ||||| ||||| |
 RESULT 8
 LOCUS B30114 623 bp DNA 13-OCT-1997
 DEFINITION T26F22.TFB TAMU Arabidopsis thaliana genomic clone T26F22, DNA sequence.
 ACCESSION B30114 GSS
 VERSION B30114
 KEYWORDS B30114.1 GI:2516080
 SOURCE GSS.
 ORGANISM thale cress.
 Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 1 (bases 1 to 623)
 Rounsley,S.D., Kelley,J.M., Field,C.E., Craven,M.B., Adams,M.D. and Venter,J.C.
 Use of a BAC End Sequence Database To Identify Minimal Overlaps for Arabidopsis Genomic Sequencing
 Unpublished (1997)
 Other_GSSs: T26F22.TFB
 Contact: Steve Rounsley
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: rounsley@tigr.org
 Seq primer: M13-21
 Class: BAC ends
 High quality sequence stop: 623.
 Location/Qualifiers
 1. 623

```

/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="T26F22"
/clone_1id="TAMU"
/sex="hermaphrodite"
/note="Vector: BelobACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"

BASE COUNT      261 a      87 c      99 g      176 t
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 623;
Best Local Similarity 65.4%; Pred. No. 8.9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Oy 5 aaaaattattccaagttctgaattttatgtttcattcttctatgttgag 64
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 378 AATATAAATTTGAAGATTTCCTTCCTGCAATTAATTTTCTCTGTTGGG 319
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Oy 65 caataagtcgaatgaag 82
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 318 TTCGAAAAAAGAAAG 301
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 9
B78178/c 627 bp DNA GSS 16-JAN-1998
LOCUS T3J11TF TAMU Arabidopsis thaliana genomic clone T3J11, DNA
DEFINITION
Sequence.
ACCESSION B78178
VERSION B78178.1 GI:2774817
KEYWORDS GSS.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 627)
Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., Adams,M.D. and Venter
,J.C.
A BAC End Sequence Database for Identifying Minimal Overlaps in
Arabidopsis Genomic Sequencing. Update 3
JOURNAL Unpublished (1997)
COMMENT Other-GSS: T3J11TR
Contact: Steve Rounsley
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: rounsley@ligr.org
Seq primer: M13-21
Class: BAC ends
High quality sequence stop: 627.
Location/Qualifiers
1. 627
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="T3J11"
/clone_1id="TAMU"
/sex="hermaphrodite"
/note="Vector: BelobACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"

BASE COUNT      261 a      87 c      100 g      178 t      1 others
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 627;
Best Local Similarity 65.4%; Pred. No. 8.9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

```

```

Oy 5 aaaaattattccaagttctgaattttatgtttcattcttctatgttgag 64
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 371 AATATAAATTTGAAGATTTCCTTCCTGCAATTAATTTTCTCTGTTGGG 312
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Oy 65 caataagtcgaatgaag 82
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 311 TTCGAAAAAAGAAAG 294
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 10
AQ958844
LOCUS AQ958844 640 bp DNA GSS 28-JAN-2000
DEFINITION LEREF16TR LERE Arabidopsis thaliana genomic clone LEREF16, DNA
sequence.
ACCESSION AQ958844
VERSION AQ958844.1 GI:6786545
KEYWORDS GSS.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 640)
Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Utterbach,T.,
Feldblyum,T., Jiang,F., Creasy,T. and Fraser,C.M.
Genomic survey sequencing of Landsberg erecta ecotype of
Arabidopsis thaliana and identification of sequence-based
polymorphisms
Unpublished (2000)
JOURNAL Contact: Xiaoying Lin
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: at@ligr.org
For additional information, see http://www.ligr.org/tdb/at.html
Similar to A. thaliana chloroplast sequence (GB:AP000423)
Seq primer: TR
Class: Shotgun.
Location/Qualifiers
1. 640
/organism="Arabidopsis thaliana"
/strain="Landsberg erecta"
/db_xref="taxon:3702"
/clone="LEREF16"
/clone_1id="LERE"
/note="Organ: Leaf; Vector: pUC19TK; Total genomic DNA was
sheared to 0.6-0.8 kbp before ligation."

BASE COUNT      181 a      99 c      83 g      277 t

Query Match      38.7%; Score 34.8; DB 13; Length 640;
Best Local Similarity 65.4%; Pred. No. 8.8e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

Oy 5 aaaaattattccaagttctgaattttatgtttcattcttctatgttgag 64
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 147 AATATAAATTTGAAGATTTCCTTCCTGCAATTAATTTTCTCTGTTGGG 206
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Oy 65 caataagtcgaatgaag 82
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 207 TTCGAAAAAAGAAAG 224
    ||| || ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 11
B98363/c 643 bp DNA GSS 31-MAR-1998
LOCUS T24C11TFB TAMU Arabidopsis thaliana genomic clone T24C11, DNA
DEFINITION
sequence.
ACCESSION B98363
VERSION B98363.1 GI:3000442

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KEYWORDS GSS:
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidops.
REFERENCE 1 (bases 1 to 643)
AUTHORS Rounsley,S.D., Field,C.E., Baas,S., Linher,K., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., Adams,M.D. and Venter,
J.C.
TITLE A BAC End Sequence Database for Identifying Minimal Overlaps in
Arabidopsis Genomic Sequencing. Update 3
JOURNAL Unpublished (1997)
COMMENT Other_GSSs: T24C1TF T24C1TRB
Contact: Steve Rounsley
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: rounsley@tigr.org
Seq primer: M13-21
Class: BAC ends
High quality sequence stop: 643.
FEATURES
source
location/Qualifiers
1. 643
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="T24C11"
/clone_lib="TAMU"
/sex="hermaphrodite"
/note="Vector: BeloBACII; Site_1: HindIII; Site_2: HindIII
; Produced by Rod Wing"
BASE COUNT 268 a 90 c 103 g 181 t 1 others
ORIGIN

Query Match 38.7%, Score 34.8; DB 13; Length 643;
Best Local Similarity 65.4%; Pred. No. 8.8e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
OY 5 aaaaattattccaagttactcgaattttatcttcacagttctctatgttgag 64
DB 378 AATATAAAATTTGAAGATTTCATTCTTCTGAATTAATTTTCTCTGTGGGG 319
OY 65 caataagtcacaatgaag 82
DB 318 TTCTGCAAAAAAAGAGAG 301

RESULT 12
CNS00WZM/C DNA GSS 28-JUN-1999
LOCUS 660 bp
DEFINITION Arabidopsis thaliana genome survey sequence T7 end of BAC T1202 of
TAMU library from strain Columbia of Arabidopsis thaliana, genomic
survey sequence.
ACCESSION AL094241
VERSION AL094241.1 GI:5295395
KEYWORDS GSS.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 660)
AUTHORS Salanoubat,M., Choise,N., Attiguenave,F., Brotlier,P., Wincker,P.,
Samson,D., Saurin,W., Weissenbach,J. and Queller,F.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 660)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (25-JUN-1999) Genoscope - Centre National de Sequencage :

```

BP 191 91006 ENVY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- web : www.genoscope.cns.fr)
FEATURES
    source
        1..660
            /organism="Arabidopsis thaliana"
            /strain="Columbia"
            /db_xref="taxon:3702"
            /clone_lib="FRMU"
            /clone="T1202"
            /note="end : T7"
BASE COUNT      272 a      93 c      107 g      188 t
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 660;
Best Local Similarity 65.4%; Pred. No. 8.7e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

OY      5 aaaaattattccaagttactgaattttatbgtttcagttttcttatgttgaag 64
        ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      371 AATATATAAATTGGAAGTTTGATTTCTTCTTCATTAATTAATTTTCTTCTGTGGGG 312
        ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
OY      65 caataagtcacatgaaag 82
        ||||| ||||| |||||
Db      311 TTCTCAAAAAAAGAAAG 294

RESULT 13
A0968517      683 bp      DNA      GSS      28-JAN-2000
LOCUS      LERJ03TR LERG Arabidopsis thaliana genomic clone LERJ03, DNA
DEFINITION      sequence.
ACCESSION      A0968517
VERSION      A0968517.1 GI:6796218
KEYWORDS      GSS.
SOURCE      thale cress.
ORGANISM      Arabidopsis thaliana
                Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
                Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
                Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE      1 (bases 1 to 683)
                Buell,C.R., Lin,X., Pal,G., Barnstead,M., Bowman,C., Uterbach,T.,
                Feldblyum,T., Liang,F., Creasy,T. and Fraser,C.M.
                Genomic survey sequencing of Landsberg erecta ecotype of
                Arabidopsis thaliana and identification of sequence-based
                polymorphisms
                Unpublished (2000)
JOURNAL      Contact: Xiaoying Lin
COMMENT      The Institute for Genomic Research
                9712 Medical Center Dr., Rockville, MD 20850, USA
                Tel: 301 838 0200
                Fax: 301 838 0208
                Email: atetlgr.org
                For additional information, see http://www.tigr.org/tdb/at.html
                Similar to A. thaliana chloroplast sequence (GB:AP000423)
                Seq primer: "TR
                Class: Shotgun.
FEATURES
    source
        1..683
            /organism="Arabidopsis thaliana"
            /strain="Landsberg erecta"
            /db_xref="taxon:3702"
            /clone="LERJ03"
            /clone_lib="LERJ03"
            /clone="T1202"
            /note="Organ: Leaf; Vector: pUC19tk; Total genomic DNA was
                sheared to 0.4-0.7 kbp before ligation."
BASE COUNT      208 a      111 c      88 g      276 t
ORIGIN

Query Match      38.7%; Score 34.8; DB 13; Length 683;
Best Local Similarity 65.4%; Pred. No. 8.5e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

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OY	5	aaaaatatttcgaagttactcgttaatttatagtttcttatgttgag	64
Db	512	AAATTAATAATTGAAGTTTGATTTCTTCCTCGAATTAATTTTTTCTCCTGTGGG	571
OY	65	caataagtcgaatgaag	82
Db	572	TTC TGAAAAAAAAGAAG	589
RESULT	14		
LOCUS	A0958265/c		
DEFINITION	LERRAW47TF LERA Arabidopsis thaliana genomic clone LERRAW47, DNA sequence.		
ACCESSION	A0958265		
VERSION	A0958265.1	GI:6785966	
KEYWORDS	GSS.		
SOURCE	Thale cress.		
ORGANISM	Arabidopsis thaliana		
REFERENCE	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.		
AUTHORS	Buell,C.R., Lin,X., Pai,G., Barnstead,M., Bowman,C., Uterbach,T., Feldblum,T., Liang,F., Creasy,T. and Fraser,C.M.		
TITLE	Genomic survey sequencing of Landsberg erecta ecotype of Arabidopsis thaliana and identification of sequence-based polymorphisms		
JOURNAL	Unpublished (2000)		
COMMENT	Contact: Xiaoying Lin The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850, USA Tel.: 301 838 0200 Fax: 301 838 0208 Email: atel@ig.org For additional information, see http://www.tigr.org/tdb/at/al.html Similar to A. thaliana chloroplast sequence (GB:AF000423) Seq primer: TF Class: shotgun.		
FEATURES	Location/Qualifiers		
source	1..743 /organism="Arabidopsis thaliana" /strain="Landsberg erecta" /db_xref="taxon:3702" /clone="LERRAW47" /clone_id="LERA" /note="Organ: Leaf; Vector: PHOSI; Total genomic DNA was sheared to 0.9-1 Kbp before ligation."		
BASE COUNT	311 a 107 c 114 g 211 t		
ORIGIN			
Query Match	38.7%; Score 34.8; DB 13; Length 743;		
Best Local Similarity	65.4%; Pred. No. 8.2e+02;		
Matches	51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;		
OY	5	aaaaatatttcgaagttactcgttaatttatagtttcttatgttgag	64
Db	534	AAATTAATAATTGAAGTTTGATTTCTTCCTCGAATTAATTTTTTCTCCTGTGGG	475
OY	65	caataagtcgaatgaag	82
Db	474	TTC TGAAAAAAAAGAAG	457
RESULT	15		
LOCUS	A0956484		
DEFINITION	LERRAK26TR LERA Arabidopsis thaliana genomic clone LERRAK26, DNA sequence.		
ACCESSION	A0956484		
VERSION	A0956484.1	GI:6784185	

KEYWORDS	GSC.
SOURCE	thale cress.
ORGANISM	Arabidopsis thaliana
REFERENCE	Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsids. 1 (bases 1 to 798)
AUTHORS	Buell,C.R., Lin,X., Pai,G., Barnstead,M., Bowman,C., Ullrich,T., Feldblum,T., Liang,F., Creasy,T. and Fraser,C.M.
TITLE	Genomic survey sequencing of landsberg erecta ecotype of Arabidopsis thaliana and identification of sequence-based polymorphisms
JOURNAL	Unpublished (2000)
COMMENT	Contact: Xiaoying Lin The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850, USA Tel.: 301 838 0200 Fax: 301 838 0208 Email: at@ligr.org For additional information, see http://www.ligr.org/tcb/at/at.html Similar to A. thaliana chloroplast sequence (CB:AT000423) Seq primer: TR Class: Shotgun.
FEATURES	Location/Qualifiers
source	1..798 /organism="Arabidopsis thaliana" /strain="landsberg erecta" /db_xref="taxon:3702" /clone="LERAK26" /clone_id="LERA"
BASE COUNT	/note="Organ: Leaf; Vector: PHOSI; Total genomic DNA was sheared to 0.9-1 Kbp before ligation." 231 a 115 c 109 g 343 t
ORIGIN	
Query Match	38.7%; Score 34.8; DB 13; Length 798;
Best Local Similarity	65.4%; Pred. No. 7.9e+02;
Matches 51; Conservative 0; Mismatches 27; Indels 0; Gaps 0;	
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Search completed: January 24, 2002, 01:51:33
Job time: 2485 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:18:46 ; Search time 90.4 Seconds
(Without alignments)

225.476 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgacaaaaattatttcaaa.....gtccaatgaagcaagtgc 90

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 351203 seqs, 113238999 residues

Total number of hits satisfying chosen parameters: 702406

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database : Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	31.2	34.7	3337	2 US-08-719-822B-1	Sequence 1, Appl
C 3	31.2	34.7	3337	4 US-09-092-458-1	Sequence 1, Appl
C 4	30.6	34.0	87350	3 US-08-781-891-79	Sequence 79, Appl
C 5	30	33.3	4140	3 US-08-884-731-2	Sequence 2, Appl
C 6	29.6	32.9	1703	4 US-08-378-313-18	Sequence 18, Appl
C 7	28.2	31.3	2334	1 US-08-062-632-4	Sequence 4, Appl
C 8	28	31.1	1013	1 US-07-920-519-30	Sequence 30, Appl
C 9	28	31.1	1013	1 US-08-086-410-23	Sequence 23, Appl
C 10	28	31.1	1013	1 US-08-314-586-30	Sequence 30, Appl
C 11	27.8	30.9	1584	1 US-07-667-376A-1	Sequence 1, Appl
C 12	27.4	30.4	1100	4 US-07-861-458C-4	Sequence 4, Appl
C 13	27.4	30.4	5613	2 US-08-463-418-1	Sequence 1, Appl
C 14	27.2	30.2	602	1 US-08-764-100-8	Sequence 8, Appl
C 15	27.2	30.2	642	1 US-08-764-100-13	Sequence 13, Appl
C 16	27.2	30.2	662	1 US-08-764-100-7	Sequence 7, Appl
C 17	27.2	30.2	662	1 US-08-998-416-185	Sequence 185, App
C 18	27.2	30.2	663	4 US-08-998-416-191	Sequence 191, App
C 19	27.2	30.2	663	4 US-08-998-416-937	Sequence 937, App
C 20	27.2	30.2	701	4 US-08-998-416-701	Sequence 701, App
C 21	27.2	30.2	711	4 US-08-998-416-786	Sequence 786, App
C 22	27.2	30.2	724	4 US-08-998-416-683	Sequence 683, App
C 23	27.2	30.2	732	4 US-08-998-416-1036	Sequence 1036, App
C 24	27.2	30.2	767	4 US-08-998-416-472	Sequence 472, App
C 25	27.2	30.2	782	4 US-08-998-416-224	Sequence 224, App
C 26	27.2	30.2	827	4 US-08-998-416-535	Sequence 535, App
C 27	27.2	30.2	828	4 US-08-998-416-538	Sequence 538, App

28	27.2	30.2	834	4 US-08-998-416-305	Sequence 305, App
C 29	27.2	30.2	854	4 US-08-998-416-534	Sequence 534, App
C 30	27.2	30.2	860	4 US-08-998-416-287	Sequence 287, App
C 31	27.2	30.2	2993	1 US-08-764-100-2	Sequence 2, Appl
C 32	27.2	30.2	2993	1 US-08-764-100-10	Sequence 10, Appl
C 33	27.2	30.2	3000	1 US-08-764-100-9	Sequence 9, Appl
C 34	27.2	30.2	3001	1 US-08-764-100-1	Sequence 1, Appl
C 35	27.2	30.2	14176	1 US-08-307-499-1	Sequence 1, Appl
C 36	27.2	30.2	14176	1 US-08-307-499-14	Sequence 14, Appl
C 37	27.2	30.2	14176	4 US-09-299-268-1	Sequence 1, Appl
C 38	27.2	30.2	14176	4 US-09-299-268-14	Sequence 14, Appl
C 39	27	30.0	642	3 US-08-817-926-49	Sequence 49, Appl
C 40	27	30.0	839	3 US-08-817-926-50	Sequence 50, Appl
C 41	27	30.0	1478	3 US-08-817-926-1	Sequence 1, Appl
C 42	27	30.0	1478	3 US-09-297-053-2	Sequence 2, Appl
C 43	27	30.0	3562	3 US-08-817-926-19	Sequence 19, Appl
C 44	26.8	29.8	2168	3 US-08-749-522-6	Sequence 6, Appl
C 45	26.8	29.8	7083	4 US-09-198-839-1	Sequence 1, Appl

ALIGNMENTS

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RESULT 1
US-08-072-610-1/c
: Sequence 1, Application US/08072610
: Patent No. 5532133
:
: GENERAL INFORMATION:
: APPLICANT: Barnwell, John
: TITLE OF INVENTION: Plasmidium vivax Blood Stage Antigens,
: TITLE OF INVENTION: Monoclonal Antibodies, and Diagnostic Assays
: NUMBER OF SEQUENCES: 2
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Darby and Darby
: STREET: 805 Third Ave.
: CITY: New York
: STATE: New York
: COUNTRY: USA
: ZIP: 10022-7513
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25
:
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/072.610
: FILING DATE: 19930602
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: Gogoris, Adga
: REGISTRATION NUMBER: 29,714
: REFERENCE/DOCKET NUMBER: 5986/07686
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (212)527-7700
: TELEFAX: (212)753-6237
: TELEX: 236687
:
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 3337 base pairs
: TYPE: NUCLEIC ACID
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHEICAL: NO
: ANTI-SENSE: NO
: ORIGINAL SOURCE:
: ORGANISM: Plasmidium vivax
: IMMEDIATE SOURCE:
: CLONE: pVMB3.3.1
:
: US-08-072-610-1

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34.7%; Score 31.2; DB 1; Length 3337;

RESULT 3
US-09-092-458-1/c

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; Patent No. 6231861
;
; GENERAL INFORMATION:
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; APPLICANT: Barnwell, John
;
; TITLE OR INVENTION: Plasma-derived human blood stage Antigen

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; TITLE OF INVENTION: Monoclonal Antibodies, and Diagnostic Assays
;
; NUMBER OF SEQUENCES: 4
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; CORRESPONDENCE ADDRESS:
;
; Address: Durham and Durham
;

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; CITY: New York
; STATE: New York
; COUNTY:

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1  COUNTRY: USA
2  ZIP: 10022-7513
3
4  COMPUTER READABLE FORM:
5
6  MEDIUM TYPE: Floppy disk
7
8  COMPUTER: IBM PC compatible
9
10 OPERATING SYSTEM: PC-DOS/MS-DOS
11
12 SOFTWARE: PatentL Release #1.0, Version #1.2
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14 CURRENT APPLICATION DATA:
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16 APPLICATION NUMBER: US/09/092,458

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; FILING DATE:
 ; CLASSIFICATION:
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 08/719,822
 ; FILING DATE: 09/30/96
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ATTORNEY/AGENT INFORMATION:
NAME: Gogoris, Adda
REGISTRATION NUMBER: 29,714
REFERENCE/DOCKET NUMBER: 5986/17666053
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)527-7700

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: TELEFAX: (212) 753-0237
:
: TELEX: 236687
:
: INFORMATION FOR SEQ ID NO: 1:
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: SEQUENCE CHARACTERISTICS:
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: LENGTH: 3337 base pairs
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: TYPE: nucleic acid
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: STRANDEDNESS: double
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;      TOPOLOGY: linear
;      MOLECULE TYPE: DNA (genomic)
;      HYPOTHETICAL: NO
;      ANTI-SENSE: NO

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ORIGINAL SOURCE:
ORGANISM: *Plasmodium vivax*

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; IMMEDIATE SOURCE:
; CLONE: PVMB3.3.1
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05-09-092-458-1

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Best Local Similarity	63.28;

Matches	48; Conservative
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[illegible]

RESULT 4
US-08-781-891-79

; Sequence 79, Application US/08781891
; Patent No. 6090620

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;; GENERAL INFORMATION:
; APPLICANT: Fu, Ying-Hui
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APPLICANT: Yu, Chang-Eu
APPLICANT: Oshima, Junko


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Query Match          34.0%; Score 30.6; DB 3; Length 87350;
Best Local Similarity 62.3%; Pred. No. 9;
Matches 48; Conservative 0; Mismatches 29; Indels 0; Gaps 0

QY      1 atgaaaaaatatttcacaaagttacgtcgtlaattttcattgatttcattgtt 60
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Db 76835 ATGCATATAGGGATATTTGTGAAGGTTCTGTGATGTTTCTGTAGAAAAGTTATCTCAAGG 76894

QY      61 ggagcaataagttccaat 77
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Db 76895 GCATATATATCTTCATTT 76911

RESULT      5
US-08-894-731-2
; Sequence 2, Application US/08894731
; Patent No. 6084089
; GENERAL INFORMATION:
; APPLICANT: MINE, Toshiki
; APPLICANT: OHYAMA, Akio
; APPLICANT: HIYOSHI, Toru
; APPLICANT: KASAKA, Keisuke
; TITLE OF INVENTION: COLD-INDUCIBLE PROMOTER SEQUENCE
; FILE REFERENCE: 760-234P
; CURRENT APPLICATION NUMBER: US/08/894,731
; CURRENT FILING DATE: 1997-10-27
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 4140
; TYPE: DNA
; ORGANISM: Solanum tuberosum
US-08-894-731-2

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RESULT 6
US-08-378-313-18
: Sequence 18, Application US/08378313
: Patent No. 6207881
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: GENERAL INFORMATION:
: APPLICANT: THEOLOGIS, ATHANASIOS
: APPLICANT: SATO, TAKAHIDO
: TITLE OF INVENTION: CONTROL OF FRUIT RIPENING THROUGH
: TITLE OF INVENTION: GENETIC CONTROL OF ACC SYNTHESIS
: NUMBER OF SEQUENCES: 34
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: MORRISON & FOERSTER
: STREET: 755 Page Mill Road
: CITY: Palo Alto
: STATE: California
: COUNTRY: USA
: ZIP: 94304-1018
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentln Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/378,313
: FILING DATE:
: CLASSIFICATION: 800
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/862,493
: FILING DATE: 02-APR-1992
: ATTORNEY/AGENT INFORMATION:
: NAME: MURASHIGE, KATE H.
: REGISTRATION NUMBER: 29,959
: REFERENCE/DOCKET NUMBER: 29190-20002.20
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 856-5600
: TELEFAX: (415) 494-0792
: TELEX: 706141
: INFORMATION FOR SEQ ID NO: 18:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1703 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 11..1489
US-08-378-313-18

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QY	61	ggagc	65
Db	176	TCTGC	172

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[illegible]

STREET: 3000 K Street, Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/314,586
FILING DATE: 28-SEP-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/659,408
FILING DATE: 25-APR-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16781/509/BDL
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)836-9300
TELEFAX: (703)683-4109
TELEX: 899149
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 1013 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
IMMEDIATE SOURCE:
CLONE: Fragment D
US-08-314-586-30

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QY      71 gtccaatgaaagaacag 86
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RESULT 11
US-07-667-276A-1
: Sequence 1, Application US/07667276A
: Patent No. 5470971
: GENERAL INFORMATION:
: APPLICANT: Kondo, Keiji
: APPLICANT: Inouye, Masayori
: TITLE OF INVENTION: STRESS-INDUCED PROTEINS, GENES CODING
: TITLE OF INVENTION: THEREFOR, TRANSFORMED CELLS OF ORGANISMS, METHODS AND
: TITLE OF INVENTION: APPLICATIONS
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Weisner & Associates
: STREET: 230 S. Fifteenth Street, Suite 500
: CITY: Philadelphia
: STATE: PA
: COUNTRY: USA
: ZIP: 19102
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS

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Search completed: January 24, 2002, 02:18:58
Job time: 3931 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:24:24 ; Search time 93.51 Seconds
(without alignments)
217.977 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 90
Sequence: 1 atgaaaaaatatttcaaa.....gtccaatgaagcaagtgc 90

Scoring table: OLIGO-NUC
Gapop 60.0, Gapext 60.0

Searched: 351203 seqs, 113238999 residues

Word size : 0

Total number of hits satisfying chosen parameters: 495388

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database : Issued Patents-NA:*

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4: /cgn2_6/ptodata/2/ina/6B.COMB.seq:*
5: /cgn2_6/ptodata/2/ina/PCrUS.COMB.seq:*
6: /cgn2_6/ptodata/2/ina/backfile1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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	2	13	14.4	30	2	US-08-629-001A-79		Sequence 79, Appli
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	4	13	14.4	38	5	PCT-US96-00547-40		Sequence 40, Appli
	5	13	14.4	39	1	US-08-105-483-168		Sequence 168, App
	6	13	14.4	39	1	US-08-709-209-168		Sequence 168, App
	7	13	14.4	39	1	US-08-303-275-56		Sequence 56, Appli
	8	13	14.4	39	1	US-08-458-101-168		Sequence 168, App
	9	12	13.3	14	1	US-08-271-880A-201		Sequence 201, App
	10	12	13.3	14	2	US-08-910-408-201		Sequence 201, App
	11	12	13.3	14	3	US-09-249-215-201		Sequence 201, App
	12	12	13.3	18	3	US-09-163-162-19		Sequence 19, Appli
	13	12	13.3	18	4	US-09-286-407-19		Sequence 19, Appli
	14	12	13.3	21	1	US-08-661-507-6		Sequence 6, Appli
	15	12	13.3	21	2	US-08-855-085-4		Sequence 4, Appli
	16	12	13.3	21	2	US-09-166-030-4		Sequence 4, Appli
	17	12	13.3	21	2	US-08-865-675-4		Sequence 4, Appli
	18	12	13.3	21	2	US-08-933-749-5		Sequence 5, Appli
	19	12	13.3	21	2	US-09-237-510-4		Sequence 4, Appli
	20	12	13.3	21	3	US-09-120-916-4		Sequence 4, Appli
	21	12	13.3	21	3	US-08-964-020-8		Sequence 8, Appli
	22	12	13.3	21	3	US-09-235-583-5		Sequence 5, Appli
	23	12	13.3	21	4	US-09-599-164-5		Sequence 5, Appli
	24	12	13.3	23	1	US-08-727-003A-10		Sequence 10, Appli
	25	12	13.3	23	3	US-08-487-799-11		Sequence 11, Appli
	26	12	13.3	24	2	US-08-210-762E-31		Sequence 31, Appli
	27	12	13.3	24	4	US-08-256-799-26		Sequence 26, Appli

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33	12	13.3	26	3	US-08-835-728D-72	Sequence 176, Appli
34	12	13.3	26	4	US-09-490-558-72	Sequence 176, Appli
35	12	13.3	26	4	US-09-490-558-176	Sequence 176, App
36	12	13.3	27	2	US-08-668-128B-6	Sequence 6, Appli
37	12	13.3	27	2	US-08-905-445-6	Sequence 6, Appli
38	12	13.3	27	3	US-08-644-116A-6	Sequence 6, Appli
39	12	13.3	28	2	US-08-771-602D-20	Sequence 20, Appli
40	12	13.3	28	2	US-08-771-602D-21	Sequence 21, Appli
41	12	13.3	30	1	US-08-619-724-3	Sequence 3, Appli
42	12	13.3	30	2	US-08-628-001A-25	Sequence 25, Appli
43	12	13.3	30	4	US-08-642-274D-104	Sequence 104, App
44	12	13.3	30	4	US-08-935-312-14	Sequence 14, Appli
45	12	13.3	33	1	US-08-341-456A-14	Sequence 14, Appli

ALIGNMENTS

RESULT 1
US-08-672-215-1/c
Sequence 1, Application US/08672215
Patent No. 6020121
GENERAL INFORMATION:
APPLICANT: Ying Bao, Amy Boggs, Pamela R. Contag,
APPLICANT: Nancy A. Federspiel, Alan Herbert,
APPLICANT: Scott J. Hecker, Francois Melouin
TITLE OF INVENTION: INHIBITORS OF REGULATORY PATHWAYS
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon 6 Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071-2066
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ for Windows 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/672.215
FILING DATE: June 25, 1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/004,626
FILING DATE: September 29, 1995
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard J.
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 219/158
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 488-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-672-215-1

Query Match 14.4%; Score 13; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 27 tgaattttatg 39
|||||
DB 20 tgaattttatg 8

RESULT 2

US-08-629-001A-79
; Sequence 79, Application US/08629001A
; Patent No. 5858661
; GENERAL INFORMATION:
; APPLICANT: Shiloh, Yosef
; TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
; NUMBER OF SEQUENCES: 139
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kohn & Associates
; STREET: 30500 No. 5858661thwestern Hwy;
; CITY: Farmington Hills
; STATE: Michigan
; COUNTRY: US
; ZIP: 48334
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/629,001A
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Kohn, Kenneth I.
; REGISTRATION NUMBER: 30,955
; REFERENCE/DOCKET NUMBER: 2290,00032
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (810) 539-5050
; TELEFAX: (810) 539-5055
; INFORMATION FOR SEQ ID NO: 79:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-629-001A-79

Query Match 14.4%; Score 13; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.9e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaatttttc 17
|||||
DB 15 AAAAAATTATTC 27

RESULT 3

US-08-642-274D-158
; Sequence 158, Application US/08642274D
; Patent No. 6200749
; GENERAL INFORMATION:
; APPLICANT: Shiloh, Yosef
; TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
; FILE REFERENCE: 229000033
; CURRENT APPLICATION NUMBER: US/08/642,274D
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 158

; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: intronic
US-08-642-274D-158

Query Match 14.4%; Score 13; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 6.9e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaatttttc 17
|||||
DB 15 aaaaatttttc 27

RESULT 4

PCT-US96-00547-40/c
; Sequence 40, Application PC/TUS9600547
; GENERAL INFORMATION:
; APPLICANT: Vitrogenetics Corporation
; TITLE OF INVENTION: RECOMBINANT POXYVIRUS-HTML, COMPOSITIONS
; NUMBER OF SEQUENCES: 56
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Curtis, Morris & Safford, P.C.
; STREET: 530 Fifth Avenue, 25th Floor
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/00547
; FILING DATE: 12-JAN-1996
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/372,664
; FILING DATE: 13-JAN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Frommer, William S.
; REGISTRATION NUMBER: 25,506
; REFERENCE/DOCKET NUMBER: 454310-2621
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 840-3333
; TELEFAX: (212) 840-0712
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 38 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
PCT-US96-00547-40

Query Match 14.4%; Score 13; DB 5; Length 38;
Best Local Similarity 100.0%; Pred. No. 6.8e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgttaattttat 38
|||||
DB 25 cgttaattttat 13

RESULT 5

US-08-105-483-168/c

Sequence 168, Application US/08105483
Patent No. 5494807
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
NUMBER OF SEQUENCES: 462
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/105,483
FILING DATE: 12-AUG-1993
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/847,951
FILING DATE: 06-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2400
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-105-483-168

Query Match 14.4%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 ctgtaattttat 38
|||||
Db 25 CTGTAATTTTAT 13

RESULT 6
US-08-709-209-168/C
Sequence 168, Application US/08709209
Patent No. 5762938
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
NUMBER OF SEQUENCES: 462
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/709,209
FILING DATE: 21-AUG-1996
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/105,483
FILING DATE: 12-AUG-1993
APPLICATION NUMBER: US 07/847,951
FILING DATE: 06-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2400
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-709-209-168

Query Match 14.4%; Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 ctgtaattttat 38
|||||
Db 25 CTGTAATTTTAT 13

RESULT 7
US-08-303-275-56/C
Sequence 56, Application US/08303275
Patent No. 576598
GENERAL INFORMATION:
APPLICANT: Paoletti, Enzo
APPLICANT: Tartaglia, James
APPLICANT: Cox, William I.
TITLE OF INVENTION: IMMUNODEFICIENCY VIRUS RECOMBINANT
NUMBER OF SEQUENCES: 205
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESS: c/o William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/303,275
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/897,382
FILING DATE: 11-JUN-1992
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2420
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712

INFORMATION FOR SEQ ID NO: 56:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-303-275-56

Query Match 14.4%: Score 13; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. NO. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgtgaattttat 38
|||||
DB 25 CGTGATTTTAT 13

RESULT 8
US-08-458-101-168/c
Sequence 168, Application US/08458101
Patent No. 5766599

GENERAL INFORMATION:
APPLICANT: Paolucci, Enzo
APPLICANT: Perkus, Marion E.
APPLICANT: Taylor, Jill
APPLICANT: Tartaglia, James
APPLICANT: No. 5766599, Elizabeth K.
APPLICANT: Riviere, Michel
APPLICANT: de Taisne, Charles
APPLICANT: Limbach, Keith J.
APPLICANT: Johnson, Gerard P.
APPLICANT: Pincus, Steven E.
APPLICANT: Cox, William I.
APPLICANT: Audonnet, Jean-Christophe Francis
APPLICANT: Getlig, Russell Robert
TITLE OF INVENTION: GENETICALLY ENGINEERED VACCINE
TITLE OF INVENTION: STRAIN
NUMBER OF SEQUENCES: 467
CORRESPONDENCE ADDRESS:
ADDRESSEE: Curtis, Morris & Safford
ADDRESSEE: C/O William S. Frommer
STREET: 530 Fifth Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/458,101
FILING DATE: 01-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Frommer, William S.
REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 45310-2740
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
INFORMATION FOR SEQ ID NO: 168:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-458-101-168

Query Match 14.4%: Score 13; DB 1; Length 39;

Best Local Similarity 100.0%; Pred. No. 6.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 cgtgaattttat 38
|||||
DB 25 CGTGATTTTAT 13

RESULT 9
US-08-271-880A-201

Sequence 201, Application US/08271880A
Patent No. 5693535

GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowrira
APPLICANT: James McSwigen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 MB
MEDIUM TYPE: Storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/271,880A
FILING DATE: July 7, 1994
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including Application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Waidburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-271-880A-201

Query Match 13.3%: Score 12; DB 1; Length 14;
Best Local Similarity 100.0%; Pred. NO. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
DB 2 CAATGAAGCAA 13

RESULT 10

US-08-910-408-201
Sequence 201, Application US/08910408
Patent No. 5972704
GENERAL INFORMATION:
APPLICANT: Kenneth G. Diaper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITTING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910.408
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/271.880
FILING DATE: July 7, 1994
APPLICATION NUMBER: 08/103.243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882.886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-910-408-201
Query Match 13.3%; Score 12; DB 2; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagcaaa 85
Db 2 CAATGAAGCAA 13
RESULT 11
US-09-249-215-201
Sequence 201, Application US/09249215
Patent No. 6159692
GENERAL INFORMATION:
APPLICANT: Kenneth G. Diaper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson

TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITTING
HUMAN IMMUNODEFICIENCY VIRUS
REPLICATION
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
STREET: Suite 4700
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM Compatible
MEDIUM TYPE: storage
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/249.215
FILING DATE: 12-Feb-1999
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/910.408
FILING DATE: <Unknown>
APPLICATION NUMBER: 08/103.243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882.886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 201:
US-09-249-215-201
Query Match 13.3%; Score 12; DB 3; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagcaaa 85
Db 2 CAATGAAGCAA 13
RESULT 12
US-09-163-162-19
Sequence 19, Application US/09163162
Patent No. 6077709
GENERAL INFORMATION:
APPLICANT: Bennett, C. Frank
APPLICANT: Ackermann, Elizabeth J.
APPLICANT: Swayze, Eric E.
APPLICANT: Cowsett, Lex M.
TITLE OF INVENTION: ANTISENSE MODULATION OF SURVIVIN EXPRESSION
FILE REFERENCE: RTS-0008
CURRENT APPLICATION NUMBER: US/09/163.162
CURRENT FILING DATE: 1998-09-29
NUMBER OF SEQ ID NOS: 47
SEQ ID NO 19
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence

FEATURE:
OTHER INFORMATION: Antisense Oligonucleotide
US-09-163-162-19

Query Match 13.3%; Score 12; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 51 tcttatgttg 62
Db 1 tcttatgttg 12

RESULT 13
US-09-286-407-19
Sequence 19, Application US/09286407A
Patent No. 6165788
GENERAL INFORMATION:
APPLICANT: Bennett, C. Frank
APPLICANT: Ackermann, Elizabeth J.
APPLICANT: Cowser, Lex M.
TITLE OF INVENTION: ANTISENSE MODULATION OF SURVIVIN EXPRESSION
FILE REFERENCE: ISPH-0349
CURRENT APPLICATION NUMBER: US/09/286,407A
CURRENT FILING DATE: 1999-04-05
NUMBER OF SEQ ID NOS: 48
SEQ ID NO 19
LENGTH: 18
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Antisense Oligonucleotide
US-09-286-407-19

Query Match 13.3%; Score 12; DB 4; Length 18;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 51 tcttatgttg 62
Db 1 tcttatgttg 12

RESULT 14
US-08-661-507-6/C
Sequence 6, Application US/08661507
Patent No. 5814490
GENERAL INFORMATION:
APPLICANT: Spears, Patricia A.
TITLE OF INVENTION: AMPLIFICATION AND DETECTION OF CHLAMYDIA
TITLE OF INVENTION: TRACHOMATIS NUCLEIC ACIDS
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
STREET: 1 Becton Drive
CITY: Franklin Lakes
STATE: NJ
COUNTRY: US
ZIP: 07417
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/661,507
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Fugit, Donna R.

REGISTRATION NUMBER: 32,135
REFERENCE/DOCKET NUMBER: P-3489
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-661-507-6

Query Match 13.3%; Score 12; DB 1; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgtt 50
Db 13 gtttcattgtt 2

RESULT 15
US-08-855-085-4/C
Sequence 4, Application US/08855085
Patent No. 5846726
GENERAL INFORMATION:
APPLICANT: Nadeau, James G.
APPLICANT: Pitner, James B.
APPLICANT: Schram, James L.
APPLICANT: Linn, Carl P.
APPLICANT: Vonk, Glenn P.
TITLE OF INVENTION: Detection of Nucleic Acids by
TITLE OF INVENTION: Fluorescence Quenching
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
STREET: 1 Becton Drive
CITY: Franklin Lakes
STATE: NJ
COUNTRY: US
ZIP: 07417
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/855,085
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Fugit, Donna R.
REGISTRATION NUMBER: 32,135
REFERENCE/DOCKET NUMBER: P-3747
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-855-085-4

Query Match 13.3%; Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgtt 50
Db 13 gtttcattgtt 2

RESULT 16

US-09-186-030-4/c
; Sequence 4, Application US/09186030
; Patent No. 5919630
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Pitner, James B.
; APPLICANT: Schram, James L.
; APPLICANT: Linn, Carl P.
; APPLICANT: Vonk, Glenn P.
; APPLICANT: Walker, George T.
; TITLE OF INVENTION: Detection of Nucleic Acids by
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/186,030
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/855,085
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3747
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-09-186-030-4

Query Match 13.3%, Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 17
US-08-865-675-4/c
; Sequence 4, Application US/08865675
; Patent No. 5928669
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Pitner, James B.
; APPLICANT: Linn, Carl P.
; APPLICANT: Schram, James L.
; APPLICANT: Vonk, Glenn P.
; TITLE OF INVENTION: DETECTION OF NUCLEIC ACIDS BY
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/865,675
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3746
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-865-675-4

Query Match 13.3%, Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 18
US-08-933-749-5/c
; Sequence 5, Application US/08933749
; Patent No. 5935791
; GENERAL INFORMATION:
; APPLICANT: Nadeau, James G.
; APPLICANT: Hsieh, Helen V.
; APPLICANT: Pitner, James B.
; APPLICANT: Linn, Carl P.
; TITLE OF INVENTION: Detection of Nucleic Acids by
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
; STREET: 1 Becton Drive
; CITY: Franklin Lakes
; STATE: NJ
; COUNTRY: US
; ZIP: 07417
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/933,749
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Fugit, Donna R.
; REGISTRATION NUMBER: 32,135
; REFERENCE/DOCKET NUMBER: P-3749
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-933-749-5

Query Match 13.3%, Score 12; DB 2; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 19
US-09-237-510-4/C

; Sequence 4, Application US/09237510
; Patent No. 5958700

; GENERAL INFORMATION:

; APPLICANT: Nadeau, James G.

; APPLICANT: Pitner, James B.

; APPLICANT: Linn, Carl P.

; APPLICANT: Schram, James L.

; TITLE OF INVENTION: DETECTION OF NUCLEIC ACIDS BY

; TITLE OF INVENTION: FLUORESCENCE QUENCHING

; NUMBER OF SEQUENCES: 9

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company

; STREET: 1 Becton Drive

; CITY: Franklin Lakes

; STATE: NJ

; COUNTRY: US

; ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/237,510

; FILING DATE:

; CLASSIFICATION:

; ATTORNEY/AGENT INFORMATION:

; NAME: Fugit, Donna R.

; REGISTRATION NUMBER: 32,135

; REFERENCE/DOCKET NUMBER: P-3746

; INFORMATION FOR SEQ ID NO: 4:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 21 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; US-09-237-510-4

Query Match 13.3%; Score 12; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 20
US-09-120-916-4/C

; Sequence 4, Application US/09120916
; Patent No. 6054279

; GENERAL INFORMATION:

; APPLICANT: Nadeau, James G.

; APPLICANT: Pitner, James B.

; APPLICANT: Schram, James L.

; APPLICANT: Linn, Carl P.

; APPLICANT: Vonk, Glenn P.

; APPLICANT: Walker, George T.

; TITLE OF INVENTION: Detection of Nucleic Acids by

; TITLE OF INVENTION: Fluorescence Quenching

; NUMBER OF SEQUENCES: 6

; CORRESPONDENCE ADDRESS:

ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
STREET: 1 Becton Drive
CITY: Franklin Lakes
STATE: NJ
COUNTRY: US

ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/120,916

; FILING DATE:

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/855,085

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Fugit, Donna R.

; REGISTRATION NUMBER: 32,135

; REFERENCE/DOCKET NUMBER: P-3747

; INFORMATION FOR SEQ ID NO: 4:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 21 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; US-09-120-916-4

Query Match 13.3%; Score 12; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 21
US-08-964-020-8/C

; Sequence 8, Application US/08964020
; Patent No. 6077669

; GENERAL INFORMATION:

; APPLICANT: Vonk, Glenn P.

; APPLICANT: Little, Michael C.

; TITLE OF INVENTION: Kit and Method for Fluorescence Based

; TITLE OF INVENTION: Detection Assay

; NUMBER OF SEQUENCES: 20

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Richard J. Rodrick - Becton, Dickinson and

; ADDRESS: Company

; STREET: 1 Becton Drive

; CITY: Franklin Lakes

; STATE: NJ

; COUNTRY: USA

; ZIP: 07417

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/964,020

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Higbet, David W.

; REGISTRATION NUMBER: 30,265

; REFERENCE/DOCKET NUMBER: P-4025

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (201) 847-5317

TELEFAX: (201) 848-9228
: INFORMATION FOR SEQ ID NO: 8:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-08-964-020-8

Query Match 13.3%; Score 12; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCATGTTT 2

RESULT 22
US-09-235-583-5/c
: Sequence 5, Application US/09235583
: Patent No. 6130047
: GENERAL INFORMATION:
: APPLICANT: Nadeau, James G.
: APPLICANT: Hsieh, Helen V.
: APPLICANT: Pitner, James B.
: APPLICANT: Lim, Carl P.
: TITLE OF INVENTION: Detection of Nucleic Acids by
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
: STREET: 1 Becton Drive
: CITY: Franklin Lakes
: STATE: NJ
: COUNTRY: US
: ZIP: 07417
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/235,583
: FILING DATE:
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Fugit, Donna R.
: REGISTRATION NUMBER: 32,135
: REFERENCE/DOCKET NUMBER: P-3749
: INFORMATION FOR SEQ ID NO: 5:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-09-235-583-5

Query Match 13.3%; Score 12; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCATGTTT 2

RESULT 23
US-09-599-164-5/c
: Sequence 5, Application US/09599164
: Patent No. 6261784

GENERAL INFORMATION:
: APPLICANT: Nadeau, James G.
: APPLICANT: Hsieh, Helen V.
: APPLICANT: Pitner, James B.
: APPLICANT: Lim, Carl P.
: TITLE OF INVENTION: Detection of Nucleic Acids by
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: R. J. Rodrick, Becton Dickinson and Company
: STREET: 1 Becton Drive
: CITY: Franklin Lakes
: STATE: NJ
: COUNTRY: US
: ZIP: 07417

COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/599,164
: FILING DATE:
: CLASSIFICATION:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US/08/933,749
: FILING DATE:
: ATTORNEY/AGENT INFORMATION:
: NAME: Fugit, Donna R.
: REGISTRATION NUMBER: 32,135
: REFERENCE/DOCKET NUMBER: P-3749
: INFORMATION FOR SEQ ID NO: 5:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 21 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-09-599-164-5

Query Match 13.3%; Score 12; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
DB 13 GTTTCATGTTT 2

RESULT 24
US-08-727-003A-10/c
: Sequence 10, Application US/08727003A
: Patent No. 5804383
: GENERAL INFORMATION:
: APPLICANT: Gruenert, Dieter, C.
: APPLICANT: Dohman, Austin F.
: TITLE OF INVENTION: A METHOD AND ASSAY FOR
: TITLE OF INVENTION: DETECTION OF THE EXPRESSION
: TITLE OF INVENTION: OF ALLELE-SPECIFIC MUTATIONS
: TITLE OF INVENTION: BY ALLELE-SPECIFIC IN SITU
: TITLE OF INVENTION: REVERSE TRANSCRIPTASE
: TITLE OF INVENTION: POLYMERASE CHAIN REACTION
: NUMBER OF SEQUENCES: 55
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: PETERS, VERNY, JONES & BIK A, L.L.P.
: STREET: 385 Sherman Avenue
: CITY: Palo Alto
: STATE: California
: COUNTRY: United States of America
: ZIP: 94306-1840
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette - 3.5 inch, 1.44 Kb storage
: COMPUTER: PC

OPERATING SYSTEM: DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA: 003A
APPLICATION NUMBER: US/08/727,003A
FILING DATE: October 8, 1996
CLASSIFICATION: 435
TITLE OF INVENTION: Molecular Clones Producing Recombinant DNA Antigens of
HANTAVIRUS
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 60/005,254
FILING DATE: October 10, 1995
ATTORNEY/AGENT INFORMATION:
NAME: Hana Vetry
REGISTRATION NUMBER: 30,518
REFERENCE/DOCKET NUMBER: 480-77
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415)324-1677
TELEFAX: (415)324-1678
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 23 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: synthetic oligonucleotide
US-08-727-003A-10

Query Match 13.3%; Score 12; DB 1; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

Oy 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 25
US-08-487-799-11/C
Sequence 11, Application US/08487799C
Patent No. 6010908
GENERAL INFORMATION:
APPLICANT: Gruenert, Dieter C.
APPLICANT: Kunzelmann, Karl
TITLE OF INVENTION: GENE THERAPY BY SMALL FRAGMENTS HOMOLOGOUS REPLACEMENT
FILE REFERENCE: 480.18-1(HV)
CURRENT APPLICATION NUMBER: US/08/487,799C
CURRENT FILING DATE: 1995-06-07
EARLIER APPLICATION NUMBER: 07/933,471
EARLIER FILING DATE: 1992-08-21
EARLIER APPLICATION NUMBER: 08/409,544
EARLIER FILING DATE: 1995-03-24
NUMBER OF SEQ ID NOS: 87
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 11
LENGTH: 23
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: synthetic
US-08-487-799-11

Query Match 13.3%; Score 12; DB 3; Length 23;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

Oy 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 26
US-08-210-762E-31

Sequence 31, Application US/08210762E
Patent No. 5837441
GENERAL INFORMATION:
APPLICANT: Hjelte, Brian
APPLICANT: Jensen, Steve
TITLE OF INVENTION: Molecular Clones Producing Recombinant DNA Antigens of
HANTAVIRUS
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 85
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hoffman, Wasson & Gittler
STREET: 2361 Jefferson Davis Highway
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 500 Kb storage
COMPUTER: Accel 486
OPERATING SYSTEM: Windows 3.1
SOFTWARE: Wordperfect 6.1 for Windows
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/210,762E
FILING DATE: 22-MAR-94
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/141,035
FILING DATE: 26-OCT-93
APPLICATION NUMBER: 08/120,096
FILING DATE: 13-SEP-93
APPLICATION NUMBER: 08/111,519
FILING DATE: 25-AUG-93
ATTORNEY/AGENT INFORMATION:
NAME: Butml, Jean A.
REGISTRATION NUMBER: 24,236
REFERENCE/DOCKET NUMBER: A4710CIP3.SU3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)415-0100
TELEFAX: (703)418-2768
INFORMATION FOR SEQ ID NO: 31:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA viral
HYPOTHETICAL: no
ANTI-SENSE: no
ORIGINAL SOURCE:
ORGANISM: Four Corners Hantavirus
INDIVIDUAL ISOLATE: 3H226
IMMEDIATE SOURCE:
LIBRARY:
CLONE:
PUBLICATION INFORMATION:
AUTHORS: Hjelte, Brian
AUTHORS: Jensen, Steven
AUTHORS: Torrez-Martinez, No. 5837441ah
AUTHORS: Yamada, Takashi
AUTHORS: No. 5837441te, Kurt
AUTHORS: Zumwalt, Ross
AUTHORS: MacInnes, Kersti
AUTHORS: Myers, Gerald
TITLE: A No. 5837441el Hantavirus Associated with an Outbreak of Fatal Respir
TITLE: Disease in the Southwestern United States: Evolutionary Relationships
TITLE: Hantaviruses-Running Title: Hantavirus-associated ARDS
JOURNAL: Journal of Virology
VOLUME: 68
PAGES: in press
DATE: 1994
RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 24
US-08-210-762E-31

Query Match 13.3%; Score 12; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

OY 21 ttatctgtaatt 32
|||||
Db 2 GTTACTGTAATT 13

RESULT 27
US-08-256-799-26
Sequence 26, Application US/08256799
Patent No. 6222094

GENERAL INFORMATION:
APPLICANT: HANSSON, Lennart
APPLICANT: STROEMOYIST, Mats
APPLICANT: BERGSTROM, Sven
APPLICANT: HERNNELL, Olle
APPLICANT: Toernell, Jan
TITLE OF INVENTION: DNA ENCODING KAPPA-CASEIN, PROCESS FOR
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK
STREET: 419 Seventh Street, N.W., Suite 300
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20004

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/256,799
FILING DATE: 06-DEC-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK 88/92
FILING DATE: 23-JAN-1992
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, Iver P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: HANSSON-1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-737-3528
TELEFAX: 202-737-3528
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-256-799-26

Query Match 13.3%; Score 12; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

OY 35 ttatgtttcat 46
|||||
Db 12 TTATGTTTCAT 23

RESULT 28
US-08-462-437-26
Sequence 26, Application US/08462437
Patent No. 6232094
GENERAL INFORMATION:
APPLICANT: HANSSON, Lennart

APPLICANT: STROEMOYIST, Mats
APPLICANT: BERGSTROM, Sven
APPLICANT: HERNNELL, Olle
APPLICANT: Toernell, Jan
TITLE OF INVENTION: DNA ENCODING KAPPA-CASEIN, PROCESS
NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK
STREET: 419 Seventh Street, N.W., Suite 300
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20004

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/462,437
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK 88/92
FILING DATE: 23-JAN-1992
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, Iver P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: HANSSON-1A
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-628-5197
TELEFAX: 202-737-3528
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-462-437-26

Query Match 13.3%; Score 12; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0;

OY 35 ttatgtttcat 46
|||||
Db 12 TTATGTTTCAT 23

RESULT 29
US-08-271-880A-140/C
Sequence 140, Application US/08271880A
Patent No. 5693535

GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chawitra
APPLICANT: James McSwigen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.

ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/271,880A
FILING DATE: July 7, 1994
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below:
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-271-880A-140

Query Match 13.3%; Score 12; DB 1; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagca 85
Db 23 CAATGAAGCAA 12

RESULT 30
US-08-910-408-140/C
Sequence 140, Application US/08910408
Patent No. 59/2704
GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,408

FILING DATE:
APPLICATION NUMBER: 08/271,880
FILING DATE: July 7, 1994
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-910-408-140

Query Match 13.3%; Score 12; DB 2; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 caatgaagca 85
Db 23 CAATGAAGCAA 12

RESULT 31
US-09-249-215-140/C
Sequence 140, Application US/09249215
Patent No. 6159692
GENERAL INFORMATION:
APPLICANT: Kenneth G. Draper
APPLICANT: Bharat Chowli
APPLICANT: James McSwiggen
APPLICANT: Dan T. Stinchcomb
APPLICANT: James D. Thompson
TITLE OF INVENTION: METHOD AND REAGENT FOR INHIBITING
TITLE OF INVENTION: HUMAN IMMUNODEFICIENCY VIRUS
NUMBER OF SEQUENCES: 232
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lyon & Lyon
STREET: 633 West Fifth Street
CITY: Los Angeles
STATE: California
COUNTRY: U.S.A.
ZIP: 90071
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
MEDIUM TYPE: storage
COMPUTER: IBM Compatible
OPERATING SYSTEM: IBM P.C. DOS 5.0
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/249,215
FILING DATE: 12-Feb-1999
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/910,408
FILING DATE: <unknown>
APPLICATION NUMBER: 08/103,243
FILING DATE: August 6, 1993
APPLICATION NUMBER: 07/882,886
FILING DATE: May 14, 1992
ATTORNEY/AGENT INFORMATION:

NAME: Warburg, Richard
REGISTRATION NUMBER: 32,327
REFERENCE/DOCKET NUMBER: 206/116
TELECOMMUNICATION INFORMATION:
TELEPHONE: (213) 489-1600
TELEFAX: (213) 955-0440
TELEX: 67-3510
INFORMATION FOR SEQ ID NO: 140:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 140:
US-09-249-215-140

Query Match 13.3%; Score 12; DB 3; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 74 caatgaagcaa 85
Db 23 CAATGAAGCAA 12

RESULT 32
US-08-835-728D-72
Sequence 72, Application US/08835728D
Patent No. 6017704
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,728D
FILING DATE: April 11, 1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/656,716
FILING DATE: June 03, 1996,
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 72:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-835-728D-72

Query Match 13.3%; Score 12; DB 3; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 32 tttaatgttt 43
Db 6 TTTTATGTTTT 17

RESULT 33
US-08-835-728D-176/C
Sequence 176, Application US/08835728D
Patent No. 6017704
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/835,728D
FILING DATE: April 11, 1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/656,716
FILING DATE: June 03, 1996,
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 176:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-835-728D-176

Query Match 13.3%; Score 12; DB 3; Length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 32 tttaatgttt 43
Db 21 TTTTATGTTTT 10

RESULT 34
US-09-490-558-72
Sequence 72, Application US/09490558
Patent No. 6265171
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylin, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400

CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/490,558
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/835,728
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 72:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-09-490-558-72

Query Match 13.3% Score 12; DB 4; length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 ttttatgttt 43
|||||
DB 6 ttttatgttt 17

RESULT 35
US-09-490-558-176/c
Sequence 176, Application US/09490558
Patent No. 6265171
GENERAL INFORMATION:
APPLICANT: Herman, James G.
APPLICANT: Baylan, Stephen B.
TITLE OF INVENTION: Methylation Specific Detection
NUMBER OF SEQUENCES: 216
CORRESPONDENCE ADDRESS:
ADDRESSER: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/490,558
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/835,728
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347

REFERENCE/DOCKET NUMBER: 07265/125001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 176:
SEQUENCE CHARACTERISTICS:
LENGTH: 26 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-09-490-558-176

Query Match 13.3% Score 12; DB 4; length 26;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 ttttatgttt 43
|||||
DB 21 ttttatgttt 10

RESULT 36
US-08-668-128B-6/c
Sequence 6, Application US/08668128B
Patent No. 5840568
GENERAL INFORMATION:
APPLICANT: Pfeundschuh, Michael
TITLE OF INVENTION: Hodgkin's Disease Associated Molecules And
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSER: Felte & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/668,128B
FILING DATE: 21-JUNE-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/644,116
FILING DATE: 10-MAY-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/580,980
FILING DATE: 03-JANUARY-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/479,328
FILING DATE: 07-JUNE-1995
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5840568man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5441
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 27 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-668-128B-6

Query Match 13.3% Score 12; DB 2; length 27;

Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
Db 26 TGAAGCAAGTG 15

RESULT 37

US-08-905-445-6/c
Sequence 6, Application US/08905445
Patent No. 5864015

GENERAL INFORMATION:

APPLICANT: Pfeundsich, Michael
TITLE OF INVENTION: Hodgkin's Disease Associated Molecules And
TITLE OF INVENTION: Uses Thereof
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/905,445

FILING DATE: 04-AUG-1997

CLASSIFICATION: 424

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/668,128

FILING DATE: 21-JUNE-1996

APPLICATION NUMBER: 08/644,116

FILING DATE: 10-MAY-1996

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/580,980

FILING DATE: 03-JANUARY-1996

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/479,328

FILING DATE: 07-JUNE-1995

ATTORNEY/AGENT INFORMATION:

NAME: Hanson, No. 5864015man D.

REGISTRATION NUMBER: 30,946

REFERENCE/DOCKET NUMBER: LUD 5441

TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 688-9200

TELEFAX: (212) 838-3884

INFORMATION FOR SEQ ID NO: 6:

SEQUENCE CHARACTERISTICS:

LENGTH: 27 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-08-905-445-6

Query Match 13.3%; Score 12; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
Db 26 TGAAGCAAGTG 15

RESULT 38

US-08-644-116A-6/c
Sequence 6, Application US/08644116A
Patent No. 6140464

GENERAL INFORMATION:

APPLICANT: Pfeundsich, Michael; Rammensee, Hans-Georg
TITLE OF INVENTION: Method For Identifying Or Isolating A Molecule
TITLE OF INVENTION: And Molecules Identified Thereby
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage

COMPUTER: IBM

OPERATING SYSTEM: PC-DOS

SOFTWARE: Wordperfect

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/644,116A

FILING DATE: 10-MAY-1996

CLASSIFICATION: 436

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/580,980

FILING DATE: 03-JANUARY-1996

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/479,328

FILING DATE: 07-JUNE-1995

ATTORNEY/AGENT INFORMATION:

NAME: Hanson, No. 6140464man D.

REGISTRATION NUMBER: 30,946

REFERENCE/DOCKET NUMBER: LUD 5410.2

TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 688-9200

TELEFAX: (212) 838-3884

INFORMATION FOR SEQ ID NO: 6:

SEQUENCE CHARACTERISTICS:

LENGTH: 27 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-08-644-116A-6

Query Match 13.3%; Score 12; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 77 tgaagcaagt 88
Db 26 TGAAGCAAGTG 15

RESULT 39

US-08-771-602D-20
Sequence 20, Application US/08771602D
Patent No. 5976795

GENERAL INFORMATION:

APPLICANT: Voytas, Daniel F.

APPLICANT: Zou, Siye

TITLE OF INVENTION: Retrotransposon and Methods

NUMBER OF SEQUENCES: 51

CORRESPONDENCE ADDRESS:

ADDRESSEE: Greenlee, Winner and Sullivan, P.C.

STREET: 5370 Manhattan Circle, Suite 201

CITY: Boulder

STATE: Colorado

COUNTRY: USA

ZIP: 80303

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/771,602D

FILING DATE: 20-DEC-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/010,869
FILING DATE: 31-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Ferber, Donna M.
REGISTRATION NUMBER: 33,878
REFERENCE/DOCKET NUMBER: 8-96
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 499-8080
TELEFAX: (303) 499-8089
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 28 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "Oligonucleotide."
HYPOTHETICAL: NO
US-08-771-602D-20

Query Match 13.3%; Score 12; DB 2; Length 28;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 31 atttattgttt 42
|||||
Db 7 ATTATTATGTTT 18

RESULT 40
US-08-771-602D-21/C
Sequence 21, Application US/08771602D
Patent No. 5976795
GENERAL INFORMATION:
APPLICANT: Voytas, Daniel F.
APPLICANT: Zou, Sige
TITLE OF INVENTION: Retrotransposon and Methods
NUMBER OF SEQUENCES: 51
CORRESPONDENCE ADDRESS:
ADDRESSEE: Greenlee, Winner and Sullivan, P.C.
STREET: 5370 Manhattan Circle, Suite 201
CITY: Boulder
STATE: Colorado
COUNTRY: USA
ZIP: 80303
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/771,602D
FILING DATE: 20-DEC-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/010,869
FILING DATE: 31-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Ferber, Donna M.
REGISTRATION NUMBER: 33,878
REFERENCE/DOCKET NUMBER: 8-96
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 499-8080
TELEFAX: (303) 499-8089
INFORMATION FOR SEQ ID NO: 21:
SEQUENCE CHARACTERISTICS:
LENGTH: 28 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "Oligonucleotide."
HYPOTHETICAL: NO
US-08-771-602D-21

Query Match 13.3%; Score 12; DB 2; Length 28;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 31 atttattgttt 42
|||||
Db 22 ATTATTATGTTT 11

RESULT 41
US-08-619-724-3/C
Sequence 3, Application US/08619724
Patent No. 5827653
GENERAL INFORMATION:
APPLICANT: SAMMES, Peter George
APPLICANT: GARMAN, Andrew John
TITLE OF INVENTION: NUCLEIC ACID DETECTION WITH ENERGY TRANSFER
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: PILLSBURY MADISON & SUTRO, L.L.P.
STREET: 1100 New York Avenue, N.W.
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20005-3918
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Microsoft Word
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/619,724
FILING DATE: 20-MAY-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/GB94/02068
FILING DATE: 23-SEP-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9412106.8
FILING DATE: 16-JUN-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: GB 9319826.5
FILING DATE: 23-SEP-1993
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-619-724-3

Query Match 13.3%; Score 12; DB 1; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 26 cgttaattttta 37
|||||
Db 29 CTTGATATTTTA 18

RESULT 42
US-08-629-001A-25/C
Sequence 25, Application US/08629001A
Patent No. 5858661
GENERAL INFORMATION:

APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
NUMBER OF INVENTION: GENOMIC ORGANIZATION
NUMBER OF SEQUENCES: 139
CORRESPONDENCE ADDRESS:
ADDRESSEE: Kohn & Associates
STREET: 30500 No. 5858661thwestern Hwy.
CITY: Farmington Hills
STATE: Michigan
COUNTRY: US
ZIP: 48334
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/629,001A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Kohn, Kenneth I.
REGISTRATION NUMBER: 30,955
REFERENCE/DOCKET NUMBER: 2290.00032
TELECOMMUNICATION INFORMATION:
TELEPHONE: (810) 539-5050
TELEFAX: (810) 539-5055
INFORMATION FOR SEQ ID NO: 25:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-629-001A-25

Query Match 13.3%; Score 12; DB 2; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 4 aaaaaattatt 15
Db 19 AAAAAATTATT 8
RESULT 43
US-08-642-274D-104/c
Sequence 104, Application US/08642274D
Patent No. 6200749
GENERAL INFORMATION:
APPLICANT: Shiloh, Yosef
TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
TITLE OF INVENTION: SCREEN FOR A PARTIAL A-T PHENOTYPE
FILE REFERENCE: 229000033
CURRENT APPLICATION NUMBER: US/08/642,274D
CURRENT FILING DATE: 1996-05-03
NUMBER OF SEQ ID NOS: 220
SOFTWARE: Patentln Ver. 2.1
SEQ ID NO 104
LENGTH: 30
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence:intronic
OTHER INFORMATION: sequence
US-08-642-274D-104

Query Match 13.3%; Score 12; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 4 aaaaaattatt 15

Db 19 AAAAAATTATT 8
RESULT 44
US-08-935-312-14/c
Sequence 14, Application US/08935312
Patent No. 6207455
GENERAL INFORMATION:
APPLICANT: CHANG, Lung-Ji
TITLE OF INVENTION: LENTIVIRAL VECTORS
NUMBER OF SEQUENCES: 20
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK, P.L.L.C.
STREET: 624 Ninth Street, N.W.
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20001
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/935,312
FILING DATE: 22-SEP-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, Iver P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: CHANG-112
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-628-5197
TELEFAX: 202-737-3528
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "DNA"
US-08-935-312-14

Query Match 13.3%; Score 12; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 74 caatgaagcaa 85
Db 28 CAATGAAGCAA 17
RESULT 45
US-08-341-456A-14/c
Sequence 14, Application US/08341456A
Patent No. 5767074
GENERAL INFORMATION:
APPLICANT: Besmer, Peter
APPLICANT: No. 5767074ka, Karl
APPLICANT: Buck, Jochen
APPLICANT: Moore, Malcolm A.S.
TITLE OF INVENTION: COMPOSITIONS OF SOLUBLE C-KIT LIGAND AND
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Cooper & Dunham LLP
STREET: 1185 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: U.S.A.

ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/341,456A
FILING DATE: 17-NOV-1994
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: White, John P.
REGISTRATION NUMBER: 28, 678
REFERENCE/DOCKET NUMBER: 37454-C1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 278-0400
TELEFAX: (212) 391-0525
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 33 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
US-08-341-456A-14

Query Match 13.3%; Score 12; DB 1; length 33;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 42 ttcatgtttct 53
|||||
Db 21 TTCATCTTTCT 10

Search completed: January 24, 2002, 03:24:26
Job time: 3713 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:57:29 ; Search time 2099.46 Seconds
(without alignments)
460.652 Million cell updates/sec

Title: US-09-531-438-4
Perfect score: 90
Sequence: 1 atgaaaaaattatttcaaa.....gtccaatgaaagcaagtgcga 90

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 11351937 seqs, 5372889281 residues

Word size: 0

Total number of hits satisfying chosen parameters: 80718

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

Database:

EST: *
1: em_estfun: *
2: em_esthum: *
3: em_estin: *
4: em_estom: *
5: em_estpl: *
6: em_estba: *
7: em_estro: *
8: em_estov: *
9: em_hic: *
10: gb_estl: *
11: gb_estl2: *
12: gb_hic: *
13: gb_gss: *
14: em_gss_fun: *
15: em_gss_hum: *
16: em_gss_inv: *
17: em_gss_pln: *
18: em_gss_pro: *
19: em_gss_rtd: *
20: em_gss_vrt: *
21: em_gss_other: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	14	15.6	50	13	TA154F10Q
2	13	14.4	25	13	AZ829725 T. brucei
3	13	14.4	26	13	AZ829725 T. brucei
4	13	14.4	28	13	AZ43522 T. brucei
5	13	14.4	36	13	AZ43522 T. brucei
6	13	14.4	49	10	AA500776 T. brucei
7	13	14.4	50	10	AA500776 T. brucei
8	12	13.3	23	13	AZ579583 T. brucei
9	12	13.3	24	13	AZ579583 T. brucei
10	12	13.3	25	13	AZ579583 T. brucei
11	12	13.3	27	10	AZ246455 T. brucei
12	12	13.3	30	13	AZ610538 T. brucei

13	12	13.3	33	13	AZ507698
14	12	13.3	38	13	AZ487251
15	12	13.3	41	13	AZ662545
16	12	13.3	42	11	T17635
17	12	13.3	45	10	AZ514394
18	12	13.3	46	10	A1805932
19	12	13.3	48	13	AZ832106
20	12	13.3	50	11	D19972
21	11	12.2	20	13	AZ782616
22	11	12.2	21	13	AZ786088
23	11	12.2	22	10	AA991150
24	11	12.2	22	13	AZ766483
25	11	12.2	23	13	AZ387817
26	11	12.2	24	13	TA242F03P
27	11	12.2	25	13	AZ462654
28	11	12.2	28	13	AZ809415
29	11	12.2	29	11	D18726
30	11	12.2	29	13	AZ580321
31	11	12.2	31	13	AZ579378
32	11	12.2	32	13	AZ331642
33	11	12.2	32	13	AZ591923
34	11	12.2	33	11	U44209
35	11	12.2	33	13	AZ423204
36	11	12.2	33	13	AZ494328
37	11	12.2	35	10	AU014463
38	11	12.2	35	11	D19895
39	11	12.2	36	10	AU111149
40	11	12.2	36	13	AZ381596
41	11	12.2	36	13	AZ807406
42	11	12.2	37	10	AA647854
43	11	12.2	37	10	A1200438
44	11	12.2	37	13	AZ806836
45	11	12.2	38	11	D21038

ALIGNMENTS

RESULT 1
TA154F10Q
LOCUS
DEFINITION
T. brucei sheared genomic DNA clone 154f10, reverse sequence,
genomic survey sequence.
ACCESSION
AL473287.1 GI:11838560
VERSION
AL473287.1
KEYWORDS
GSS.
SOURCE
Trypanosoma brucei.
ORGANISM
Trypanosoma brucei.
Eukaryota; Euzoaria; Kinetoplastida; Trypanosomatidae;

REFERENCE
AUTHORS
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajendram, M.A. and Barrell, B.G.

TITLE
JOURNAL
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
nh@sanger.ac.uk

COMMENT

Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared
to give a tight size distribution (4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
Barrell, Oxford University Press, 1999).

Email: neilsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available
at <http://www.sanger.ac.uk/projects/T-brucei/>.

FEATURES

source
1..50

/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="154f10"

BASE COUNT 20 a 7 c 9 g 14 t

Query Match 15.6%; Score 14; DB 13; Length 50;

Best Local Similarity 100.0%; Pred. No. 2.6e+04;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 atcaaaaaaatatc 14
|||||
Db 10 ATGAAAAAATATAT 23

RESULT 2
AZ829725 25 bp DNA GSS 20-FEB-2001

LOCUS 2M010712AF Mouse 10kb plasmid UUGC1M library Mus musculus genomic

DEFINITION clone UUGC2M0107124 F, DNA sequence.

ACCESSION AZ829725

VERSION AZ829725.1 GI:12999549

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

TITLE 1 (bases 1 to 25)

JOURNAL Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,

COMMENT Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly

Mouse whole genome scaffolding with paired end reads from 10kb

Plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0107 row: I column: 24

Seq primer: CGTGTAAACGACGCCACT

Class: plasmid ends

High quality sequence stop: 25.

Location/Qualifiers

1..25

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC2M0107124"

/clone.lib="Mouse 10kb plasmid UUGC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of PWD42 (g114732114|gb|AF129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 14 a 3 c 4 g 4 t

Query Match 14.4%; Score 13; DB 13; Length 25;

Best Local Similarity 100.0%; Pred. No. 9.7e+04;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 34 ttatgtttatc 46
|||||
Db 24 TTTATGTTTCAAT 12

RESULT 3
TA123B12Q/c 26 bp DNA GSS 13-DEC-2000

DEFINITION T. brucei sheared genomic DNA clone 123b12, reverse sequence.

ACCESSION AL463522

VERSION AL463522.1 GI:11834032

KEYWORDS GSS.

SOURCE Trypanosoma brucei.

ORGANISM Trypanosoma brucei

REFERENCE Eukaryota; Eumetazoa; Kinetoplastida; Trypanosomatidae;

AUTHORS Trypanosoma.

TITLE 1 (bases 1 to 26)

JOURNAL Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,

COMMENT Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,

Meiville, S.E., Rajandream, M.A. and Barrell, B.G.

Direct Submission

Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing

Project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,

Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and

nhlesanger.ac.uk

Constructed at the Institute for Genomic Research (TIGR),

Rockville, MD. Genomic DNA isolated from a cloned population of

Trypanosoma brucei (TREU927/4 G9rat 10.1) was mechanically sheared

to give a tight size distribution (4 kb). The v + i method used for the library construction is

described in detail in Smith, H. and Venter, J.C. (Making small

insert libraries for whole genome shotgun sequencing projects. In

Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.

Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org

Details of T. brucei sequencing at the Sanger Centre are available

at http://www.sanger.ac.uk/projects/T-brucei/

Location/Qualifiers

1..26

/organism="Trypanosoma brucei"

/strain="TREU927"

/db_xref="taxon:5691"

/clone="123b12"

BASE COUNT 12 a 0 c 0 g 14 t

ORIGIN

Query Match 14.4%; Score 13; DB 13; Length 26;

Best Local Similarity 100.0%; Pred. No. 9.5e+04;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaaatattc 16
|||||

Db 20 AAAAAAATATATT 8

RESULT 4

AZ452653 28 bp DNA GSS 04-OCT-2000

DEFINITION 1M0252E07R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 ACCESSION AZ452653
 VERSION AZ452653.1 GI:10609676
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murine; Mus.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel.: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0252 row: E column: 07
 Seq primer: CACACAGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 28.
 Location/Qualifiers
 1..28
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0252E07"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g11473214(gb)AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."
 BASE COUNT 14 a 2 c 2 g 10 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 13; Length 28;
 Best Local Similarity 100.0%; Pred. No. 9.3e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 4 aaaaaattattt 16
 ||||||||||||
 Db 16 AAAAAATTATT 28
 RESULT 5
 AZ314238

LOCUS AZ314238 36 bp DNA 29-SEP-2000
 DEFINITION 1M0030N24R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 ACCESSION AZ314238
 VERSION AZ314238.1 GI:10359929
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murine; Mus.
 REFERENCE 1 (bases 1 to 36)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel.: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0030 row: N column: 24
 Seq primer: CACACAGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 36.
 Location/Qualifiers
 1..36
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0030N24"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of PMD42 (g11473214(gb)AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."
 BASE COUNT 8 a 3 c 5 g 20 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 13; Length 36;
 Best Local Similarity 100.0%; Pred. No. 8.5e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 45 atgtttcttatt 57
 ||||||||||||
 Db 2 ATGTTTCTTATT 14
 RESULT 6
 AZ314238

AA500776 49 bp mRNA EST 01-JUL-1997
 LOCUS AA500776
 DEFINITION yg01b11.1 Soares mouse NbMH Mus musculus cDNA clone IMAGE:66061
 5 Similar to TR:G133638 G133638 PARADOXASE 2. ; mRNA sequence.
 ACCESSION AA500776
 VERSION AA500776.1 GI:2235743
 KEYWORDS EST.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 49)
 REFERENCE 1 Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
 Giesel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
 Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
 Waterston,R.
 TITLE The WashU-HHMI Mouse EST Project
 JOURNAL Unpublished (1996)
 COMMENT Contact: Marra M/Mouse EST Project
 WashU-HHMI Mouse EST Project
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 MG1:504149
 Trace considered overall poor quality
 Possible reversed clone; similarity on wrong strand
 Seq primer: -28m13 rev2 ET from Amersham
 High quality sequence stop: 1.
 FEATURES
 source location/Qualifiers
 1..49
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="IMAGE:860061"
 /clone_lib="Soares mouse NbMH"
 /sex="male"
 /tissue_type="heart"
 /dev_stage="4 weeks"
 /lab_host="DH10B"
 /note="Vector: pT73D-Pac (Pharmacia) with a modified
 polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
 was primed with a Not I - oligo(dT) primer [5'
 TGTATCCATCTGAGTGGAGCGCCGGAAGTTTGTGTGTGTGTGTGTGTGT
 3']; double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not
 I and Eco RI sites of the modified pT73 vector. RNA
 provided by Dr. Minoru Ko, Wayne State Univ. Library
 constructed and normalized by Bento Soares and M.Fatima
 Bonaldo."
 BASE COUNT 17 a 8 c 10 g 14 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 10; Length 49;
 Best Local Similarity 100.0%; Pred. No. 7.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 30 aattttatgttt 42
 |||||||
 Db 25 AATTTTATGTTT 37
 RESULT 7
 LOCUS AU102750 50 bp mRNA EST 05-APR-2001
 DEFINITION AU102750 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
 CAS10730, mRNA sequence.
 ACCESSION AU102750

VERSION AU102750.1 GI:13552271
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 50)
 REFERENCE 1 Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata
 H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo
 K., Suyama,A. and Sugano,S.
 TITLE Fine structural analysis of transcription start sites of human
 JOURNAL mRNAs using full-length enriched and 5'-end enriched cDNA libraries
 COMMENT Unpublished (2001)
 Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: yusuzuki@ims.u-tokyo.ac.jp
 Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
 S. Construction and characterization of a full length-enriched and
 a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
 FEATURES
 source location/Qualifiers
 1..50
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="CAS10730"
 /clone_lib="Sugano Homo sapiens cDNA library"
 BASE COUNT 19 a 13 c 8 g 10 t
 ORIGIN
 Query Match 14.4%; Score 13; DB 10; Length 50;
 Best Local Similarity 100.0%; Pred. No. 7.6e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 73 ccaatgaagca 85
 |||||||
 Db 8 CCAATGAAGCAA 20
 RESULT 8
 LOCUS A2579583/C 23 bp DNA GSS 13-DEC-2000
 DEFINITION IM0367N03F Mouse 10kb plasmid UGCGM library Mus musculus genomic
 clone UGCGM0367N03 F, DNA sequence.
 ACCESSION A2579583
 VERSION A2579583.1 GI:11694012
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 23)
 REFERENCE 1 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenan,E., Pedersen,T., Reilly
 M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 JOURNAL plasmid inserts
 COMMENT Unpublished (2000)
 Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0367 row: N column: 03
 Seq primer: CGTGTAAACGACGCGCCAGT
 Class: plasmid ends
 High quality sequence stop: 23.

FEATURES
source

Location/Qualifiers

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1. .23
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M0367N03"
/clone_lib="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g11473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

BASE COUNT
ORIGIN

```
9 a 2 c 0 g 12 t
```

Query Match 13.3%; Score 12; DB 13; Length 23;

Best Local Similarity 100.0%; Pred. No. 2.9e+05;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 aaaaattatt 15

Db 15 AAAAAATTATT 4

RESULT 9
A2996858/c

LOCUS A2996858 24 bp DNA GSS 27-APR-2001

DEFINITION 2M0283L08F Mouse 10kb plasmid U08C2M library Mus musculus genomic

clone U08C2M0283L08 F, DNA sequence.

ACCESSION A2996858

VERSION A2996858.1 GI:13868085

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus.

REFERENCE 1 (bases 1 to 24)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

UNPUBLISHED (2000)

CONTACT: Robert B. Weiss

UNIVERSITY OF UTAH

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0283 row: L column: 08

Seq primer: CGTGTAAACGACGCCACGT

Class: plasmid ends

FEATURES
sourceHigh quality sequence stop: 24.
Location/Qualifiers

```
1. .24
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C2M0283L08"
/clone_lib="Mouse 10kb plasmid U08C2M library"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (female) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g11473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

BASE COUNT
ORIGIN

```
12 a 4 c 3 g 5 t
```

Query Match 13.3%; Score 12; DB 13; Length 24;

Best Local Similarity 100.0%; Pred. No. 2.8e+05;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 33 ttatatgttttc 44

Db 21 TTTATGTTTTC 10

RESULT 10

LOCUS A2345616 25 bp DNA GSS 29-SEP-2000

DEFINITION 1M0080E20F Mouse 10kb plasmid U08C1M library Mus musculus genomic

clone U08C1M0080E20 F, DNA sequence.

ACCESSION A2345616

VERSION A2345616.1 GI:10424853

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM Mus musculus.

REFERENCE 1 (bases 1 to 25)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

UNPUBLISHED (2000)

CONTACT: Robert B. Weiss

UNIVERSITY OF UTAH

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0080 row: E column: 20

Seq primer: CGTGTAAACGACGCCACGT

Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers

1. 25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0080E20"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (g1473214|9b|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
16 a 1 c 0 g 8 t

Query Match 13.3%; Score 12; DB 13; Length 25;
Best Local Similarity 100.0%; Pred. No. 2.8e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 29 taattttttgt 40
|||||

RESULT 11

AM246455

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AW246455. 27 bp mRNA EST 07-JAN-2000
2821693.3prime NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2821693 3',
mRNA sequence.
AW246455
AW246455.1 GI:6589448
EST.
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 27)
NIH-MGC http://mgs.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Other_ESTS: 2821693.5prime
Contact: Robert Strausberg, Ph.D.
Email: cgaabs-r@mail.nih.gov
Tissue procurement: DCTD/DPF CDNA Library Preparation: Ling
Hong/Rubin Laboratory CDNA Library Arrayed by: The I.M.A.G.E.
Consortium (LNU) DNA Sequencing by: Berkeley MGC sequencing
project Clome distribution: MGC clone distribution information can
be found through the I.M.A.G.E. Consortium/LNU at:
www.hio.lnl.gov/bbrp/image/image.html Base Calling / Quality
Scores: PHRED from University of Washington Genome Center. Vector
Trimming: cross_match from University of Washington Genome Center
PHRAP suite. Poly-T Identification: patmatch.pl from Berkeley
Drosophila Genome Project. University of Washington Genome Center:
http://www.genome.washington.edu Low Quality Sequence: 10

contiguous PHRED high quality bases following vector sequence. Very
Low Quality Sequence: Trace file contained 27 contiguous distinct
peaks following vector sequence. Polyadenylation: Based upon the
presence of a XhoI site followed by a run of 14 or more T residues
at the beginning of the sequence, this cDNA insert was
polyadenylated.
Plate: LICM7 row: H column: 14
High quality sequence stop: 10.
Location/Qualifiers

1. 27
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2821693"
/clone_lib="NIH_MGC_7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/note="Organ: lung; Vector: pORF7; Site_1: XhoI; Site_2:
ECORI; cDNA made by oligo-dT priming. Directionally
cloned into ECORI/XhoI sites using the following 5'
adaptor: GGCACGAG(C). Size selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the Laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT
ORIGIN
2 a 1 c 2 g 22 t

Query Match 13.3%; Score 12; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 2.7e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 40 ttttcagtttc 51
|||||

RESULT 12

A2610538

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

A2610538 30 bp DNA GSS 13-DEC-2000
1M0435E23R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
clone UUCG1M0435E23 R, DNA sequence.
A2610538
A2610538.1 GI:11732728
GSS.
house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 30)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0435 row: E column: 23
Seq primer: CACACAGCAACACCTATGACC
Class: plasmid ends
High quality sequence stop: 30.
Location/Qualifiers
1. 30

```
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0435E23"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-."
/notes="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g11473211419b/AP129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

```
source
1. 33
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0349021"
/clone.lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-."
/notes="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g11473211419b/AP129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."
```

```
Query Match 13.3%; Score 12; DB 13; Length 30;
Best Local Similarity 100.0%; Pred. No. 2.6e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 27 tgtattttttat 38
|||||
Db 4 TGTAAATTTTAT 15
```

```
Query Match 13.3%; Score 12; DB 13; Length 33;
Best Local Similarity 100.0%; Pred. No. 2.5e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 40 ttccaatgtttt 51
|||||
Db 16 TTTTCATGTTT 27
```

```
RESULT 13
AZ507698 33 bp DNA GSS 05-OCT-2000
LOCUS
DEFINITION
IM0349021F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0349021 F, DNA sequence.
ACCESSION
AZ507698
VERSION
AZ507698.1 GI:10689014
KEYWORDS
GSS.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
```

```
RESULT 14
AZ487251 38 bp DNA GSS 05-OCT-2000
LOCUS
DEFINITION
IM0316A18R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0316A18 R, DNA sequence.
ACCESSION
AZ487251
VERSION
AZ487251.1 GI:10654814
KEYWORDS
GSS.
SOURCE
house mouse.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
```

```
REFERENCE
AUTHORS
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
```

```
REFERENCE
AUTHORS
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
```

```
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
```

```
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
```

```
TITLE
Unpublished (2000)
JOURNAL
COMMENT
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SIC, UT
84112, USA
```

```
TITLE
Unpublished (2000)
JOURNAL
COMMENT
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SIC, UT
84112, USA
```

```
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Seg primer: 0349 row: 0 column: 21
Class: plasmid ends
High quality sequence stop: 33.
Location/Qualifiers
```

```
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0316 row: A column: 18
Seg primer: CACACAGAAACAGCATGACG
Class: plasmid ends
High quality sequence stop: 38.
```

FEATURES
source
Location/Qualifiers
1. 38
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0316A18"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gb|AF129072.1), a copy number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
14 a 14 c 5 g 5 t

Query Match
Best Local Similarity 13.3%; Score 12; DB 13; Length 38;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 Tgaataaacta 13
|||||
Db 26 TCAAAAAAATTA 37

RESULT 15
AZ662545 41 bp DNA GSS 14-DEC-2000
LOCUS 1M0541P07R Mouse 10kb plasmid UUCG1M library Mus musculus genomic
DEFINITION clone UUCG1M0541P07 R, DNA sequence.
ACCESSION AZ662545
VERSION AZ662545.1 GI:11799691
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 41)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tinney, A., von Niederhausern, A. and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0541 row: P column: 07
Seq primer: CACACAGGAACAGCTATGACC
Class: plasmid ends

JOURNAL
COMMENT

FEATURES
source
High quality sequence stop: 41.
Location/Qualifiers
1. 41
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0541P07"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gb|AF129072.1), a copy number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
16 a 9 c 5 g 11 t

Query Match
Best Local Similarity 13.3%; Score 12; DB 13; Length 41;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 52 ctatattgtga 63
|||||
Db 35 CTTATTGTTGGA 24

RESULT 16
T17635 42 bp mRNA EST 06-JUN-1994
LOCUS mps v360 The blue guys library Saccharomyces cerevisiae cDNA
DEFINITION sequence upstream of lacZ fusion similar to GAC1, X63941, mRNA sequence.
ACCESSION T17635
VERSION T17635.1 GI:459560
KEYWORDS EST.
SOURCE baker's yeast.
ORGANISM Saccharomyces cerevisiae
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes; Saccharomycetales; Saccharomycetaceae; Saccharomyces. 1 (bases 1 to 42)
Burns, N., Grimwade, B., Ross-Macdonald, P. B., Choi, E.-Y., Finberg, K., Roeder, G. S. and Snyder, M.
Large-scale analysis of gene expression, protein localization and gene disruption in Saccharomyces cerevisiae
Genes Dev. 8, 1087-1105 (1994)
95011603
Contact: Snyder M
Department of Biology
Yale University
New Haven CT 06520-8103
Tel: 2034326139
Fax: 2034326161
Email: snymp@yalevm.ycc.yale.edu
LacZ fusion; Vegetative expression; Beta-gal fusion localization pattern;
50 cytoplasmic spots; Disruption phenotype: none detected; Fusion: codon 407 of GAC1 gene. Sequence below near or adjacent to lacZ.

JOURNAL
COMMENT

FEATURES
source

Seq primer: lacZ sequences in transposon.
Location/Qualifiers
1. 42
/organism="Saccharomyces cerevisiae"
/db_xref="taxon:4932"
/clone_id="The blue guys library"
/lab_host="E.coli"
/note="Vector: PRECMTn: A yeast genomic DNA library was prepared in the vector pHS6, and subjected to transposon mutagenesis with mtn3. This mini-transposon carries lacZ sequences that lack an initiation codon; expression of lacZ is only provided by in frame fusion to yeast coding sequence. The yeast genomic DNA carrying the transposon was excised from pHS6 and transplanted back onto the yeast chromosome. Yeast colonies expressing lacZ were screened for in a color assay. A plasmid containing the genomic DNA/lacZ fusion junction was recovered from each individual yeast colony that expressed lacZ activity. These recovered plasmids comprise 'The blue guys library'. The fusion junction was then sequenced to identify the expressed ORF upstream of the fusion."

BASE COUNT
ORIGIN
7 a 12 c 11 g 12 t

Query Match 13.3%; Score 12; DB 11; Length 42;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 aatgaagcaag 86
|||||
Db 22 AATGAAGCAAG 11

RESULT 17
AZ514394

LOCUS 45 bp DNA GSS 05-OCT-2000
DEFINITION 1M0361J02F Mouse 10kb plasmid UNGC1M library Mus musculus genomic
clone UNGC1M0361J02 F, DNA sequence.

ACCESSION AZ514394
VERSION AZ514394.1 GI:10695710
KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciuromorphi; Muridae; Murinae; Mus.
1 (bases 1 to 45)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

TITLE Unpublished (2000)

JOURNAL Contact: Robert B. Weiss
COMMENT University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA

FEATURES
source
1. 45
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UNG1M0361J02"

/clone_id="Mouse 10kb plasmid UNGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42ny. Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1147321141gb1AF129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
24 a 2 c 9 g 10 t

Query Match 13.3%; Score 12; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 aaaaaaatatt 15
|||||
Db 3 AAAAAATTATT 14

RESULT 18
A1805932

LOCUS 46 bp mRNA EST 13-DEC-1999
DEFINITION t622903.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:2090356 3 similar to TR:Q99766 Q99766 HYPOTHETICAL 15.7 KD
PROTEIN: ; mRNA sequence.

ACCESSION A1805932
VERSION A1805932.1 GI:5392498
KEYWORDS EST.

SOURCE human.
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 46)
NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

TITLE Unpublished (1997)

JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgapbs-remail.nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 606 Std Error: 0.00
Seq primer: -40UP from Gldco
High quality sequence stop: 1.

FEATURES
source
1. 46
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2090356"
/clone_id="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"

/note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site:1: Not I; Site:2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (fetal lung NBHL19W, testis NHT, and B-cell NCI-CCAP-GCB1) were mixed, and ss circles were made in

vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 297480-302087, 682632-687239, 726408-728711, and 729096-731399. Subtraction by Bento Soares and M. Fatima Bonaldo.

BASE COUNT 13 a 9 c 5 g 19 t
ORIGIN

Query Match 13.3%; Score 12; DB 10; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 25 actgaattttt 36
DB 34 ACTGTAATTTT 45

RESULT 19
AZ832106/c

LOCUS 48 bp DNA GSS 20-FEB-2001
DEFINITION 2M0112113F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0112113 F, DNA sequence.

ACCESSION AZ832106
VERSION AZ832106.1 GI:13002014

KEYWORDS GSS.

SOURCE house mouse.
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 48)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0112 row: 1 column: 13
Seq primer: CGTTGTAACGACGCGCAGT

Class: plasmid ends
High quality sequence stop: 48.

FEATURES
SOURCE location/Qualifiers
1. 48

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0112113"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative of PMD42 (gi14732114|gb1AE129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 11 c 3 g 20 t
ORIGIN

Query Match 13.3%; Score 12; DB 13; Length 48;
Best Local Similarity 100.0%; Pred. No. 2.2e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
DB 48 AAAAAATTATT 37

RESULT 20
D19972/c

LOCUS 50 bp mRNA EST 30-JUL-1996
DEFINITION HUMGS00937 Human promyelocyte Homo sapiens cDNA clone mm06d08 3', mRNA sequence.

ACCESSION D19972
VERSION D19972.1 GI:500869

KEYWORDS EST.
SOURCE human.

ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo. 1 (bases 1 to 50)

AUTHORS Okubo,K., Fukushima,A., Yoshii,J., Niyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing

TITLE Unpublished (1993)

JOURNAL

COMMENT Contact: Okubo,K., Fukushima,A., Yoshii,J., Niyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.

FEATURES
SOURCE location/Qualifiers
1. 50

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="mm06d08"
/clone_lib="Human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type = promyelocyte."

BASE COUNT 20 a 5 c 11 g 14 t
ORIGIN

Query Match 13.3%; Score 12; DB 11; Length 50;
Best Local Similarity 100.0%; Pred. No. 2.2e+05;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 ttcaaatgttac 26
DB 20 TTCAAGTTTAC 9

RESULT 21
AZ782616

LOCUS 20 bp DNA GSS 16-FEB-2001
DEFINITION 2M0023F17R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC2M0023F17 R, DNA sequence.

ACCESSION AZ782616
VERSION AZ782616.1 GI:12916517

KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 20)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0023 row: F column: 17
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 20.
Location/Qualifiers
1. .20
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG2M0023F17"
/clone_lib="Mouse 10kb plasmid UUCGCM library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gb|AF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT 6 a 4 c 1 g 9 t
ORIGIN
Query Match 12.2%; Score 11; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.8e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 33 ttttatgtttt 43
|||||
Db 9 ttttatgtttt 19
RESULT 22
A2768088 21 bp DNA GSS 16-FEB-2001
LOCUS A2768088
DEFINITION IM0567G21R Mouse 10kb plasmid UUCGCM library Mus musculus genomic clone UUCGCM0567G21 R, DNA sequence.
ACCESSION A2768088

VERSION A2768088.1 GI:12886839
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus.
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 21)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0567 row: G column: 21
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 21.
Location/Qualifiers
1. .21
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCGCM0567G21"
/clone_lib="Mouse 10kb plasmid UUCGCM library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114[gb|AF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT 5 a 3 c 1 g 12 t
ORIGIN
Query Match 12.2%; Score 11; DB 13; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 40 tttcatgttt 50
|||||
Db 11 tttcatgtttt 21
RESULT 23
AA991150 22 bp mRNA EST 03-JUN-1998
LOCUS AA991150/c
DEFINITION os40a07.s1 NCI_CGAP_Br2 Homo sapiens cDNA IMAGE:1607796 3' similar to TR:Q35990 Q35990 HYPOTHETICAL 8.9 KD PROTEIN.; contains

element 11 repetitive element ;, mRNA sequence.
 AA991150
 VERSION AA991150.1 GI:317639
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 1 (bases 1 to 22)
 NCI-CCAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CCAP),
 Tumor Gene Index
 Unpublished (1997)
 JOURNAL Contact: Robert Strausberg, Ph.D.
 COMMENT Email: cgapbs-rt@mail.nih.gov
 Tissue Procurement: Christopher Moskalko, M.D., Ph.D., Michael R.
 Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CCAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbtp/image/image.html

Trace considered overall poor quality
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 1.
 Location/Qualifiers
 1..22
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1607796"
 /clone_lib="NCI-CCAP Br2"
 /sex="female, pooled"
 /tissue_type="breast"
 /lab_host="DH10B"
 /note="Vector: pRT3D-Pac (Pharmacia) with a modified
 polylinker; 1st strand cDNA was prepared from pooled bulk
 breast tumor tissue, and was then primed with a Not I -
 oligo(dT) primer. Double-stranded cDNA was ligated to Eco
 RI adaptors (Pharmacia), digested with Not I and cloned
 into the Not I and Eco RI sites of the modified pRT3
 vector. This library is the normalized version of
 NCI-CCAP-Br1.1. Library was constructed by Bento Soares
 and M. Fatima Bonaldo."

BASE COUNT 16 a 3 c 2 g 1 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 22;
 Best Local Similarity 100.0%; Pred. No. 8.5e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 46 tgtttcttat 56
 |||||||||
 Db 22 TGTTCCTTAT 12

RESULT 24
 A2766483 22 bp DNA GSS 16-FEB-2001
 LOCUS 1M056410F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 DEFINITION clone UUGC1M056410 F, DNA sequence.
 ACCESSION A2766483
 VERSION A2766483.1 GI:12883604
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 22)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly

, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhause, R.
 and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 1000 Std Error: 0.00
 Plate: 0564 row: A column: 10
 Seg primer: CGTGTAAACGACGCCAGT
 Class: plasmid ends
 High quality sequence stop: 22.
 Location/Qualifiers
 1..22
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M056410"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. coli strain XL10-Gold, Tl-resistant, F-"
 /note="Vector: pMD42nv: Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (<http://www.jax.org/resources/documents/dnares/>). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD42 (91473211419b1AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT 4 a 2 c 4 g 12 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 22;
 Best Local Similarity 100.0%; Pred. No. 8.5e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 46 tgtttcttat 56
 |||||||||
 Db 4 TGTTCCTTAT 14

RESULT 25
 A2387817 23 bp DNA GSS 02-OCT-2000
 LOCUS 1M0147824R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 DEFINITION clone UUGC1M0147824 R, DNA sequence.
 ACCESSION A2387817
 VERSION A2387817.1 GI:10501525
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 23)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,

Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL
COMMENT
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0147 row: B column: 24
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 23.

FEATURES
SOURCE
Location/Qualifiers

1. .23
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U08C1M0147B24"
/clone_lib="Mouse 10kb plasmid U08C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g1147321149b1Af129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
5 a 4 c 3 g 11 t
Query Match 12.2%; Score 11; DB 13; Length 23;
Best Local Similarity 100.0%; Pred. No. 8.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Gy 28 gtaattttat 38
|||||
Db 6 gtaattttat 16

RESULT 26
TA242F03P 24 bp DNA GSS 13-DEC-2000
LOCUS
DEFINITION T. brucei sheared genomic DNA clone 242F03, forward sequence, genomic survey sequence.
ACCESSION
VERSION AL482984
KEYWORDS AL482984.1 GI:11848725
SOURCE GSS.
ORGANISM Trypanosoma brucei.
Trypanosoma brucei
Eukaryota, Euglenozoa, Kinetoplastida, Trypanosomatidae;
Trypanosoma.
REFERENCE 1 (bases 1 to 24)
Location/Qualifiers

AUTHORS

Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.E., Rajandream, M.A. and Barrall, B.G.

TITLE

JOURNAL
COMMENT
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrall@sanger.ac.uk and nh@sanger.ac.uk

COMMENT

Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 G09at 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrall, Oxford University Press, 1999).

Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available at <http://www.sanger.ac.uk/projects/T-brucei/>.

FEATURES
SOURCE
Location/Qualifiers

1. .24
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="242f03"
BASE COUNT
ORIGIN
1 a 3 c 10 g 10 t
Query Match 12.2%; Score 11; DB 13; Length 24;
Best Local Similarity 100.0%; Pred. No. 8.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Gy 52 ctattgttg 62
|||||
Db 8 cttattgttg 18

RESULT 27
LOCUS
DEFINITION A2462654 25 bp DNA GSS 04-OCT-2000
1M0269P09R Mouse 10kb plasmid U08C1M library Mus musculus genomic clone U08C1M0269P09 R, DNA sequence.
ACCESSION
VERSION A2462654
KEYWORDS A2462654.1 GI:10620695
SOURCE GSS.
ORGANISM Mus musculus.
house mouse.

REFERENCE
AUTHORS
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.

TITLE

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL

COMMENT
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunne@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0269 row: P column: 09
Seq primer: CACACAGCAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 25.
Location/Qualifiers

FEATURES

source

1. 25
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C1M0269P09"
/clone_lib="Mouse 10Kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
10 a 2 c 0 g 13 t

Query Match
Best Local Similarity 12.2%; Score 11; DB 13; Length 25;
Matches 11: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 6 aaaaattattt 16
|||||
Db 4 AAAAATTATT 14

RESULT 28
A2809415/C 28 bp DNA GSS 20-FEB-2001
LOCUS 2M0073120F Mouse 10kb plasmid U06C1M library Mus musculus genomic
DEFINITION clone U06C2M0073120 F, DNA sequence.
ACCESSION A2809415
VERSION A2809415.1 GI:12975693
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 28)
REFERENCE
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingle,A., von Niederhausen,A.
and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10Kb
Plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0073 row: 1 column: 20
Seq primer: CGTTTAACGACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 28.

FEATURES
source

Location/Qualifiers
1. 28
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="U06C2M0073120"
/clone_lib="Mouse 10Kb plasmid U06C1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PMD42nv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g11473211419b1AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN
7 a 5 c 5 g 11 t

Query Match
Best Local Similarity 12.2%; Score 11; DB 13; Length 28;
Matches 11: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 74 caatgaagca 84
|||||
Db 14 CAATGAAGCA 4

RESULT 29
D18726 29 bp mRNA EST 12-DEC-1995
LOCUS MUSGS01788 Mouse 3'-directed Mus musculus domesticus cDNA clone
DEFINITION md0169 3', mRNA sequence.
ACCESSION D18726
VERSION D18726.1 GI:1100695
KEYWORDS EST.
SOURCE western European house mouse.
ORGANISM Mus musculus domesticus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 29)
REFERENCE
AUTHORS Kawamoto,S., Okubo,K., Yoshii,J., Katsuki,M. and Matsubara,K.
TITLE Analysis of gene expression in mouse embryogenesis by 3'-directed
cDNA sequencing
JOURNAL Unpublished (1995)
COMMENT Contact: Kawamoto,S., Okubo,K., Yoshii,J., Katsuki,M. and Matsubara,
K.
Institute for Cellular and Molecular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
FEATURES
source
Location/Qualifiers
1. 29
/organism="Mus musculus domesticus"
/strain="C57BL/6J"
/db_xref="taxon:10092"
/clone="md0169"
/clone_lib="Mouse 3'-directed"
/tissue_type="decidual tissue (day 6.5-8.5 of gestation)"
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ORIGIN
15 a 3 c 2 g 9 t

Query Match 12.2%; Score 11; DB 11; Length 29;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 36 tatgttttcac 46
 |||||||
 Db 15 TATGTTTCAT 25

RESULT 30

AZ580321 29 bp DNA GSS 13-DEC-2000
 LOCUS AZ580321
 DEFINITION IM0366G02R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0366G02 R, DNA sequence.
 ACCESSION AZ580321
 VERSION AZ580321.1 GI:11694750
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 29)
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 'M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

TITLE Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0368 row: G column: 02
 Seq primer: CACACAGAAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 29.

FEATURES
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 1..29
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 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0366G02"
 /clone_1lb="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD2 (gii4732114[gb|AF129072.1], a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT

10 a 11 c 3 g 5 t

ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 29;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 76 atgaagcaag 86
 |||||||
 Db 16 ATGAAGCAAG 26

RESULT 31

AZ579378 31 bp DNA GSS 13-DEC-2000
 LOCUS AZ579378
 DEFINITION IM0363N13R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0363N13 R, DNA sequence.
 ACCESSION AZ579378
 VERSION AZ579378.1 GI:11693807
 KEYWORDS GSS.
 SOURCE house mouse.
 ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 31)
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 'M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

TITLE Unpublished (2000)
 JOURNAL Contact: Robert B. Weiss
 COMMENT University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunne@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0363 row: N column: 13
 Seq primer: CACACAGAAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 31.

FEATURES
 SOURCE Location/Qualifiers
 1..31
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0363N13"
 /clone_1lb="Mouse 10kb plasmid UUGC1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD2 (gii4732114[gb|AF129072.1], a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

```

BASE COUNT      12 a      1 c      8 g      10 t
ORIGIN

Query Match      12.2%; Score 11; DB 13; Length 31;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      4 aaaaaaatat 14
        |||||||
DB      3 AAAAAATTTAT 13

RESULT 32
AZ331642/c      32 bp      DNA      GSS      29-SEP-2000
LOCUS
DEFINITION      1M0059P1R Mouse 10kb plasmid UGCC1M library Mus musculus genomic
ACCESSION      AZ331642
VERSION      AZ331642.1 GI:10394528
KEYWORDS
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
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Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0059 row: P column: 11
Seq primer: CACACGAGAAACAGCTAGACC
Class: plasmid ends
High quality sequence stop: 32.
Location/Qualifiers
1..32
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UCC1M0059P11"
/clone.lib="Mouse 10kb plasmid UGCC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (911473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells

```

```

BASE COUNT      14 a      7 c      1 g      10 t
ORIGIN

Query Match      12.2%; Score 11; DB 13; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      13 attcaagtt 23
        |||||||
DB      20 ATTTCAAGTT 10

RESULT 33
AZ591923      32 bp      DNA      GSS      13-DEC-2000
LOCUS
DEFINITION      1M0402F21F Mouse 10kb plasmid UGCC1M library Mus musculus genomic
ACCESSION      AZ591923
VERSION      AZ591923.1 GI:11714113
KEYWORDS
SOURCE      house mouse.
ORGANISM      Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 32)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0402 row: F column: 21
Seq primer: GGTGTAAACGACGCCACGT
Class: plasmid ends
High quality sequence stop: 32.
Location/Qualifiers
1..32
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UCC1M0402F21"
/clone.lib="Mouse 10kb plasmid UGCC1M library"
/sex="Male"
/lab.host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (911473211419b1AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into

```


Chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

Query Match 12.2%; Score 11; DB 13; Length 32;
Best Local Similarity 100.0%; Pred. No. 7.5e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 18 aaagtactg 28
Db 4 AAAGTTTACTG 14

RESULT 34
LOCUS U44209/c 33 bp mRNA EST 03-APR-1996
DEFINITION ENU44209 Aspergillus nidulans cleistothecium Emeritella nidulans
cDNA clone SE0393, mRNA sequence.

ACCESSION U44209.1 GI:1244872
VERSION U44209.1
KEYWORDS EST.
SOURCE Emeritella nidulans.
ORGANISM Emeritella nidulans.
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;

REFERENCE 1 (bases 1 to 33)
AUTHORS Lee,D., Lee,S., Hwang,H., Kim,J. and Chae,K.
TITLE Quantitative analysis of gene expression in sexual structures of
Aspergillus nidulans by sequencing of 3'-directed cDNA clones
JOURNAL FEMS Microbiol. Lett. 138 (1), 71-76 (1996)
MEDLINE 96236220
COMMENT Contact: Keon-Sang Chae
Chonbuk National University
Chonju, 561-756, S. Korea
Tel: +82-652-70-3340
Fax: +82-652-70-3345
Email: chaek@chonbuk.ac.kr.

FEATURES
SOURCE Location/Qualifiers
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/organism="Emeritella nidulans"
/strain="FGSC4"
/db_xref="taxon:162425"
/clone="SE0393"
/clone_lib="Aspergillus nidulans cleistothecium"
/tissue_type="cleistothecium"
/cell_type="Hull cell"
/dev_stage="sexual"
/note="3'-directed cDNA clones; single-pass sequencing"

BASE COUNT 18 a 3 c 2 g 10 t
ORIGIN

Query Match 12.2%; Score 11; DB 11; Length 33;
Best Local Similarity 100.0%; Pred. No. 7.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 32 ttctatgttc 42
Db 31 TTTTATGTTT 21

RESULT 35
LOCUS AZ423204 33 bp DNA GSS 03-OCT-2000
DEFINITION c1one UUGC1M0202010 F, DNA sequence.
ACCESSION AZ423204.1 GI:10547217
VERSION AZ423204.1
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 33)
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0202 row: 0 column: 10
Seq primer: CGTTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 33.
Location/Qualifiers
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/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0202010"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42mv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g114732114[gb|AF129072.1]), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent *E. coli* XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

FEATURES
SOURCE

BASE COUNT 15 a 3 c 7 g 8 t
ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 33;
Best Local Similarity 100.0%; Pred. No. 7.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 aaaaaattat 14
Db 21 AAAAAATTAT 31

RESULT 36
LOCUS AZ494328 33 bp DNA GSS 05-OCT-2000
DEFINITION 1M0329F05R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0329F05 R, DNA sequence.
ACCESSION AZ494328
VERSION AZ494328.1 GI:10668799
KEYWORDS GSS.
SOURCE house mouse.

ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 33)
 AUTHORS Dun, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C., Islam, H., Longacre, S., Mahmoud, M., Meenan, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tinney, A., von Niederhausen, A. and Wright, D., Weiss, R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT 84112 USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert length: 10000 Std Error: 0.00
 Plate: 0329 row: F column: 05
 Seq primer: CACACAGCAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 33.
 Location/Qualifiers
 1..33
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="U08C1M0329F05"
 /clone_lib="Mouse 10kb plasmid U08C1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv: Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g14732114(gblAF129072.1)), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 8 a 1 c 7 g 17 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 33;
 Best Local Similarity 100.0%; Pred. No. 7.4e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 49 ttctctatgt 59
 |||||
 Db 23 ttctctatgt 33

RESULT 37
 AU014463 35 bp mRNA EST 03-AUG-1998
 LOCUS AU014463 Schizosaccharomyces pombe late log phase cDNA
 DEFINITION AU014463 Schizosaccharomyces pombe cDNA clone spc09897, mRNA sequence.
 ACCESSION AU014463
 VERSION AU014463.1 GI:3369254
 KEYWORDS EST.

SOURCE fission yeast.
 ORGANISM Schizosaccharomyces pombe
 Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes; Schizosaccharomycetales; Schizosaccharomycetaceae; Schizosaccharomycetes.
 REFERENCE 1 (bases 1 to 35)
 AUTHORS Moriyama, M. and Mita, K.
 TITLE Identification of expressed sequence tags of Schizosaccharomyces pombe
 JOURNAL Unpublished (1998)
 COMMENT Contact: Mitsuoki Moriyama
 Genome Research Group
 National Institute of Radiological Sciences
 9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
 Email: moriyoma@nirs.go.jp
 FEATURES
 source
 1..35
 /organism="Schizosaccharomyces pombe"
 /strain="972"
 /db_xref="taxon:4896"
 /clone="spc09897"
 /clone_lib="Schizosaccharomyces pombe late log phase cDNA"
 /sex="h minus"
 /note="Vector: M13mp19: The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, http://www.nirs.go.jp)"

BASE COUNT 21 a 1 c 5 g 8 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 35;
 Best Local Similarity 100.0%; Pred. No. 7.2e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 gaaanaaata 13
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 Db 16 GAAAAAATTA 26

RESULT 38
 D19995 35 bp mRNA EST 30-JUL-1996
 LOCUS D19995 Human promyelocyte Homo sapiens cDNA clone mp1322 3', mRNA sequence.
 DEFINITION H08GS00963 Human promyelocyte Homo sapiens cDNA clone mp1322 3', mRNA sequence.
 ACCESSION D19995
 VERSION D19995.1 GI:500892
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 35)
 AUTHORS Okubo, K., Fukushima, A., Yoshii, J., Niijima, T., Kojima, Y., Yoshinari, H., Arimoto, J. and Matsubara, K.
 TITLE Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing
 JOURNAL Unpublished (1993)
 COMMENT Contact: Okubo, K., Fukushima, A., Yoshii, J., Niijima, T., Kojima, Y., Yoshinari, H., Arimoto, J. and Matsubara, K.
 Institute for Molecular and Cellular Biology
 Osaka University
 3-1 Yamada-oka, Suita, Osaka 565, Japan.
 FEATURES
 source
 1..35
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="mp1322"
 /clone_lib="Human promyelocyte"
 /note="Female, adult, cell_line = HL60, cell_type =

BASE COUNT 14 a 7 c 2 g 12 t
ORIGIN

Query Match 12.2%; Score 11; DB 11; Length 35;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 27 tgaattttta 37
|||||
Db 20 TGTAAATTTTVA 10

RESULT 39
AU111149/c 36 bp mRNA EST 19-OCT-2000
LOCUS AU111149 unpublished oligo-capped cDNA library Caenorhabditis
DEFINITION elegans cDNA yk724el 5', mRNA sequence.
ACCESSION AU111149
VERSION AU111149.1 GI:10924716
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea
; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 36)
REFERENCE Kohara, Y., Shin-i, T., Thierry-Mieg, J., Thierry-Mieg, D., Suzuki, Y.
AUTHORS A complete view of the C. elegans genome
JOURNAL Unpublished (2000)
COMMENT Contact: Yuji Kohara
Genome Biology Lab.
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.

FEATURES
SOURCE
1. 36
/organism="Caenorhabditis elegans"
/strain="N2"
/db_xref="taxon:6239"
/clone="YK724el"
/sex="Hermaphrodite"
/tissue_type="whole animal"
/dev_stage="varied"

BASE COUNT 17 a 3 c 6 g 10 t
ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 36;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 aaaaattattt 16
|||||
Db 23 AAAAATTTT 13

RESULT 40
AZ381596/c 36 bp DNA GSS 02-OCT-2000
LOCUS AZ381596
DEFINITION IM0138C16F Mouse 10kb plasmid UUC1M library Mus musculus genomic
clone UUC1M0138C16 F, DNA sequence.
ACCESSION AZ381596
VERSION AZ381596.1 GI:10495296
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 36)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0138 row: C column: 16
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 36.

FEATURES
SOURCE
1. 36
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC1M0138C16"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (g114732114[9b]AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 19 a 2 c 10 g 5 t
ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 36;
Best Local Similarity 100.0%; Pred. No. 7.2e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 44 catgtttctt 54
|||||
Db 36 CATGTTTCTT 26

RESULT 41
AZ807406/c 36 bp DNA GSS 20-FEB-2001
LOCUS AZ807406
DEFINITION 2M0070D15F Mouse 10kb plasmid UUC1M library Mus musculus genomic
clone UUC2M0070D15 F, DNA sequence.
ACCESSION AZ807406
VERSION AZ807406.1 GI:12971722
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (bases 1 to 36)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duvall,B., Hamill,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausen,A., and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10Kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT 84112, USA
 Tel.: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0070 row: D column: 15
 Seq primer: CGTGTAAACGACGCGCAGT
 Class: plasmid ends
 High quality sequence stop: 36.
 Location/Qualifiers
 1..36
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUCG2M0070D15"
 /clone_lib="Mouse 10Kb plasmid UUCG1M library"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /note="Vector: PMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PMD42 (g114732114|gblAF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 18 a 7 c 7 g 4 t
 ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 36;
 Best Local Similarity 100.0%; Pred. No. 7.2e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 26 ctgtattttt 36
 |||||||
 Db 19 CTGTATTTT 9

RESULT 42
 AA647854 37 bp mRNA EST 28-OCT-1997
 LOCUS AA647854
 DEFINITION vg080e05.s1 Knowles Solter mouse 2 cell Mus musculus cDNA clone IMAGE:1108640 5' similar to TR:G1136390 G1136390 KIA0164 PROTEIN.
 ; mRNA sequence.
 ACCESSION AA647854
 VERSION AA647854.1 GI:2574283
 KEYWORDS EST.
 SOURCE house mouse.

ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B., Theising,B., Wyllie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.
 TITLE The WashU-HHMI Mouse EST Project
 JOURNAL Unpublished (1996)
 COMMENT Contact: Marra M/Mouse EST Project
 WashU-HHMI Mouse EST Project
 Washington University School of Medicine
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through INM; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
 MGI:606808
 Trace considered overall poor quality
 Possible reversed clone; similarity on wrong strand
 High quality sequence stop: 1.
 Location/Qualifiers
 1..37
 /organism="Mus musculus"
 /strain="B6D2 F1/3"
 /db_xref="taxon:10090"
 /clone="IMAGE:1108640"
 /clone_lib="Knowles Solter mouse 2 cell"
 /tissue_type="embryo"
 /dev_stage="2-cell"
 /lab_host="DH10B"
 /note="Organ: embryo; Vector: Bluescribe (modified); site_1: MluI; site_2: SalI; Cloned unidirectionally from mRNA prepared from 13,500 2-cell stage embryos. Primer: SalI(dry): 5'-CGGCGACGCGACGCTTTT-3'. CDNAS were cloned into the MluI/SalI sites of a modified Bluescribe vector using commercial linkers (NEB). Average insert size: 1.2 kb."

BASE COUNT 9 a 8 c 8 g 12 t
 ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 37;
 Best Local Similarity 100.0%; Pred. No. 7.1e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 66 aataagtcac 76
 |||||||
 Db 26 AATAAGTCCAA 36

RESULT 43
 AI200438 37 bp mRNA EST 29-NOV-1998
 LOCUS AI200438
 DEFINITION q193b01.x1 Soares-placenta_8to9weeks_2NBHP6C09M Homo sapiens cDNA clone IMAGE:1757545 3' similar to SW:Q0PT.HUMAN Q16769 GLUTAMINYL-PEPTIDE CYCLOTRANSFERASE PRECURSOR; contains element MER35 repetitive element;; mRNA sequence.
 ACCESSION AI200438
 VERSION AI200438.1 GI:3753044
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.
Email: ccaps-remail.nih.gov
This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 863 Std Error: 0.00
Seq primer: -40UP from GIBCO
High quality sequence stop: 1.
Location/Qualifiers

FEATURES
source
1. .37
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1757545"
/clone_lib="Sceres.placenta.8to9weeks_2nbHpt09W"
/dex_stage="two placentae: one from 8 weeks and another from 9 weeks post conception"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: placenta; Vector: pT73D (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer (5' TGTTACCAATCTGAAGTGGAGGCGCGCATTTTCTTTTCTTTT 3'), double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library constructed by Bento Soares and M.Fatima Donaldso."

BASE COUNT 11 a 6 c 12 g 8 t

ORIGIN

Query Match 12.2%; Score 11; DB 10; Length 37;
Best Local Similarity 100.0%; Pred. No. 7.1e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 tgaaaaaatt 12
|||||
Db 14 TGAATAAAATT 24

RESULT 44
A2806836 37 bp DNA GSS 20-FEB-2001
LOCUS 2M0069108F Mouse 10kb plasmid UGCM1 library Mus musculus genomic
DEFINITION clone UUGC2M0069108 F, DNA sequence.
ACCESSION A2806836
VERSION A2806836.1 GI:12970584
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murineae; Mus.
1 (bases 1 to 37)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
JOURNAL Contact: Robert B. Weiss
COMMENT University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0069 row: I column: 08
Seq primer: CGTGTAAACGACGCGCACG
Class: plasmid ends
High quality sequence stop: 37.
Location/Qualifiers

FEATURES

source
1. .37
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0069108"
/clone_lib="Mouse 10kb plasmid UGCM1 library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
/note="Vector: pMD22ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD22 (gii4732114/gbiAFT29072.1) a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 14 a 1 c 10 g 12 t

ORIGIN

Query Match 12.2%; Score 11; DB 13; Length 37;
Best Local Similarity 100.0%; Pred. No. 7.1e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 30 aattttatgt 40
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Db 5 AATTTTATGT 15

RESULT 45
D21038 38 bp mRNA EST 30-JUL-1996
LOCUS HMG502021 Human promyelocyte Homo sapiens cDNA clone mp014 3',
DEFINITION mRNA sequence.
ACCESSION D21038
VERSION D21038.1 GI:504858
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 38)
Okubo,K., Fukushima,A., Yoshii,J., Niiyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
Gene expression of human promyelocytic cell line HL60 before and after induction of differentiation. A new application of 3'directed cDNA sequencing
Unpublished (1993)
JOURNAL Contact: Okubo,K., Fukushima,A., Yoshii,J., Niiyama,T., Kojima,Y., Yoshinari,H., Arimoto,J. and Matsubara,K.
COMMENT Institute for Molecular and Cellular Biology
Osaka University
3-1 Yamada-oka, Suita, Osaka 565, Japan.
Location/Qualifiers

FEATURES
source
1. .38
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="mp014"
/clone_lib="human promyelocyte"
/note="Female, adult, cell_line = HL60, cell_type = promyelocyte."
Location/Qualifiers

BASE COUNT 15 a 4 c 8 g 11 t

ORIGIN

Query Match 12.2%; Score 11; DB 11; Length 38;
 Best Local Similarity 100.0%; Pred. No. 7e+05;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 11 ttattcaag 21
 11 |||||
 Db 14 TTRTTCAAG 4

Search completed: January 24, 2002, 02:57:32
 Job time: 3954 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 03:22:36 ; Search time 1494.92 Seconds
(without alignments)
993.194 Million cell updates/sec

Title: US-09-531-438-4

Perfect score: 1 atgaaaaaattatttcaaa.....gtccaatgaagcaagtga 90

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 1472140 seqs, 8248589755 residues

Word size : 0

Total number of hits satisfying chosen parameters: 541028

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_hlg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vl.*
15: gb_vl.*
16: em_ba.*
17: em_fun.*
18: em_hum.*
19: em_in.*
20: em_om.*
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22: em_ov.*
23: em_pat.*
24: em_ph.*
25: em_pl.*
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27: em_sts.*
28: em_sy.*
29: em_un.*
30: em_vl.*
31: em_hlgo_hum.*
32: em_hlgo_inv.*
33: em_hlgo_rtd.*
34: em_hlg_hum.*
35: em_hlg_inv.*
36: em_hlg_rtd.*
36: em_hlg_other.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	13	14.4	24	6	AX164353	AX164353 Sequence
2	13	14.4	26	6	AX039624	AX039624 Sequence
3	13	14.4	26	6	AX039654	AX039654 Sequence
4	13	14.4	30	6	AR028230	AR028230 Sequence
5	13	14.4	30	6	AR138633	AR138633 Sequence
6	13	14.4	33	5	XELARSE59	K01606 Xenopus lae
7	13	14.4	36	6	AX167671	AX167671 Sequence
8	13	14.4	39	6	AR011299	AR011299 Sequence
9	13	14.4	39	6	II17937	II17937 Sequence
10	12	13.3	14	6	AR082357	AR082357 Sequence
11	12	13.3	14	6	AR120899	AR120899 Sequence
12	12	13.3	14	6	178403	178403 Sequence
13	12	13.3	16	6	AI14957	AI14957 synthetic o
14	12	13.3	18	6	AI14956	AI14956 synthetic o
15	12	13.3	18	6	AR099365	AR099365 Sequence
16	12	13.3	20	6	A64909	A64909 Synthetic a
17	12	13.3	20	6	AR05597	AR05597 Sequence
18	12	13.3	21	6	AR043539	AR043539 Sequence
19	12	13.3	21	6	AR063857	AR063857 Sequence
20	12	13.3	21	6	AR075821	AR075821 Sequence
21	12	13.3	21	6	AR096733	AR096733 Sequence
22	12	13.3	21	6	AR112325	AR112325 Sequence
23	12	13.3	21	6	E30454	E30454 Method for
24	12	13.3	21	6	E32364	E32364 Method for
25	12	13.3	21	6	E33635	E33635 Detection o
26	12	13.3	21	6	E35692	E35692 Detection o
27	12	13.3	23	6	AR037890	AR037890 Sequence
28	12	13.3	24	6	AR054522	AR054522 Sequence
29	12	13.3	24	6	AR151501	AR151501 Sequence
30	12	13.3	25	6	AR082296	AR082296 Sequence
31	12	13.3	25	6	AR120838	AR120838 Sequence
32	12	13.3	25	6	AX115268	AX115268 Sequence
33	12	13.3	25	6	178342	178342 Sequence
34	12	13.3	27	6	AR060384	AR060384 Sequence
35	12	13.3	27	6	AR117878	AR117878 Sequence
36	12	13.3	27	6	AX114027	AX114027 Sequence
37	12	13.3	28	6	AR082955	AR082955 Sequence
38	12	13.3	28	6	AR082956	AR082956 Sequence
39	12	13.3	29	6	AX155921	AX155921 Sequence
40	12	13.3	30	6	A43687	A43687 Sequence
41	12	13.3	30	6	AR028176	AR028176 Sequence
42	12	13.3	30	6	AR050260	AR050260 Sequence
43	12	13.3	30	6	AR138579	AR138579 Sequence
44	12	13.3	30	6	AR140326	AR140326 Sequence
45	12	13.3	30	6	AX020976	AX020976 Sequence

ALIGNMENTS

RESULT 1
LOCUS AX164353 24 bp DNA
DEFINITION Sequence 183 from Patent WO0138564.
ACCESSION AX164353
VERSION AX164353.1 GI:14545287
KEYWORDS
SOURCE
ORGANISM
synthetic construct.
synthetic construct
artificial sequence.
REFERENCE
1 (bases 1 to 24)
Rouleau,G.A., Latremiere,R.G., Rochefort,D., Cossette,P. and
Ragsdale,D.
Locl for idiopathic generalized epilepsy, mutations thereof and
method using same to assess, diagnose, prognosis or treat epilepsy
Patent: WO 0138564-A 183 31-MAY-2001;
JOURNAL
McGill University (CA)
FEATURES
location/Qualifiers
1..24
/organism="synthetic construct"

/db_xref="taxon:32630"
BASE COUNT 2 a 5 c 3 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 34 ttatgtttcat 46
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Db 8 TTTATGTTTCAT 20

RESULT 2
AX039624 26 bp DNA PAT 18-NOV-2000
LOCUS Sequence 13 from Patent WO0063441.
ACCESSION AX039624
VERSION AX039624.1 GI:11229653
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herinstdt,C. and Davis,R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
MITOKOR (US)

FEATURES
Source 1. 26
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"
BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 atgtttcatgtt 49
|||||
Db 7 ATGTTTCATGTT 19

RESULT 3
AX039654 26 bp DNA PAT 18-NOV-2000
LOCUS Sequence 43 from Patent WO0063441.
ACCESSION AX039654
VERSION AX039654.1 GI:11229683
KEYWORDS
SOURCE synthetic construct.
ORGANISM artificial sequence.
REFERENCE 1 (bases 1 to 26)
AUTHORS Herinstdt,C. and Davis,R.E.
TITLE Single nucleotide polymorphisms in mitochondrial genes that segregate with alzheimer's disease
JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
MITOKOR (US)

FEATURES
Source 1. 26
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR primer"
BASE COUNT 8 a 2 c 5 g 11 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 atgtttcatgtt 49
|||||
Db 7 ATGTTTCATGTT 19

RESULT 4
AR028230 30 bp DNA PAT 29-SEP-1999
LOCUS Sequence 79 from patent US 5858661.
ACCESSION AR028230
VERSION AR028230.1 GI:5940203
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shihon,Y.
TITLE Ataxia-telangiectasia gene and its genomic organization
JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
FEATURES Location/Qualifiers
Source 1. 30
/organism="unknown"
BASE COUNT 13 a 2 c 1 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaattatttc 17
|||||
Db 15 AAAAATTATTTC 27

RESULT 5
ARI38633 30 bp DNA PAT 16-JUN-2001
LOCUS Sequence 158 from patent US 6200749.
ACCESSION ARI38633
VERSION ARI38633.1 GI:14480978
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 30)
AUTHORS Shihon,Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to screen for a partial A-T phenotype
JOURNAL Patent: US 6200749-A 158 13-MAR-2001;
FEATURES Location/Qualifiers
Source 1. 30
/organism="unknown"
BASE COUNT 13 a 2 c 1 g 14 t
ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 aaaaattatttc 17
|||||
Db 15 AAAAATTATTTC 27

RESULT 6
XELARSE59

LOCUS XELARSE59 33 bp DNA VRT 28-APR-1993
 DEFINITION Xenopus laevis autonomous replication sequence e59.
 ACCESSION K01606
 VERSION K01606.1 GI:213953
 KEYWORDS autonomous replication; mutational analysis.
 SOURCE Xenopus laevis DNA.
 ORGANISM Xenopus laevis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
 Xenopodinae; Xenopus.
 REFERENCE 1 (bases 1 to 33)
 AUTHORS Keasey, S.
 TITLE Structural requirements for the function of a yeast chromosomal replicator
 JOURNAL Cell 37, 299-307 (1984)
 MEDLINE 84205653
 FEATURES
 source Location/Qualifiers
 1..33
 /organism="Xenopus laevis"
 /db_xref="taxon:8355"
 BASE COUNT 8 a 4 c 2 g 19 t
 ORIGIN

Query Match 14.4%; Score 13; DB 5; Length 33;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 31 atttattgtttt 43
 |||||||||
 1 ATTATTATGTTT 13

RESULT 7
 AXI67671 36 bp DNA PAT 03-JUL-2001
 LOCUS AXI67671
 DEFINITION Sequence 16 from Patent WO0144277.
 ACCESSION AXI67671
 VERSION AXI67671.1 GI:14597058
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM synthetic construct.
 REFERENCE 1 (bases 1 to 36)
 AUTHORS Wegrich Glover, L., Budziszewski, G.J., Levin, J.Z. and Zhou, Q.
 TITLE Herbicide target genes and methods
 JOURNAL Patent: WO 0144277-A 16 21-JUN-2001;
 Syngenta Participations AG (CH)
 FEATURES
 source Location/Qualifiers
 1..36
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="Oligonucleotide"
 BASE COUNT 6 a 4 c 11 g 15 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 36;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 36 tatgtttcatgt 48
 |||||||||
 21 TATGTTTTCATGT 33

RESULT 8
 AR011299 39 bp DNA PAT 04-DEC-1998
 LOCUS AR011299/c
 DEFINITION Sequence 168 from patent US 5762938.
 ACCESSION AR011299
 VERSION AR011299.1 GI:3969289
 KEYWORDS

SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 39)
 AUTHORS Paolletti, E., Perkus, M.E., Taylor, J., Tartaglia, J., Norton, E.K., Riviere, M., de Taisne, C., Limbach, K.J., Johnson, G.P., Pincus, S.E., Cox, M.I., Audonnet, J.F., Francis and Gettig, R. Robert.
 TITLE Modified recombinant vaccinia virus and expression vectors thereof
 JOURNAL Patent: US 5762938-A 168 09-JUN-1998;
 FEATURES
 source Location/Qualifiers
 1..39
 /organism="unknown"
 BASE COUNT 15 a 6 c 7 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
 |||||||||
 25 CTGTAATTTTAT 13

RESULT 9
 I17937 39 bp DNA PAT 07-OCT-1996
 LOCUS I17937/c
 DEFINITION Sequence 168 from patent US 5494807.
 ACCESSION I17937
 VERSION I17937.1 GI:1598292
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 39)
 AUTHORS Paolletti, E., Perkus, M.E., Taylor, J., Tartaglia, J., Norton, E.K., Riviere, M., de Taisne, C., Limbach, K.J., Johnson, G.P., Pincus, S.E., Cox, M.I., Audonnet, J.F. and Gettig, R.R.
 TITLE NYVAC Vaccinia virus recombinants comprising heterologous inserts
 JOURNAL Patent: US 5494807-A 168 27-FEB-1996;
 FEATURES
 source Location/Qualifiers
 1..39
 /organism="unknown"
 BASE COUNT 15 a 6 c 7 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 39;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ctgtaattttat 38
 |||||||||
 25 CTGTAATTTTAT 13

RESULT 10
 AR082357 14 bp DNA PAT 31-AUG-2000
 LOCUS AR082357
 DEFINITION Sequence 201 from patent US 5972704.
 ACCESSION AR082357
 VERSION AR082357.1 GI:10009083
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 14)
 AUTHORS Draper, K.G., Chowrira, B., McSwiggen, J., Stinchcomb, D.T. and Thompson, J.D.
 TITLE HIV nef targeted ribozymes
 JOURNAL Patent: US 5972704-A 201 26-OCT-1999;
 FEATURES Location/Qualifiers

source 1.14
/organism="unknown"
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 11
LOCUS AR120899 14 bp DNA PAT 16-MAY-2001
DEFINITION Sequence 201 from patent US 6159692.
ACCESSION AR120899
VERSION AR120899.1 GI:14104475
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 14)
AUTHORS Draper,K.G., Chowrira,B., McSwiggen,J., Stinchcomb,D.T. and
TITLE Thompson,J.D.
METHOD Method and reagent for inhibiting human immunodeficiency virus
replication
JOURNAL Patent: US 6159692-A 201 12-DEC-2000;
FEATURES Location/Qualifiers
source 1.14
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 12
LOCUS I78403 14 bp DNA PAT 03-APR-1998
DEFINITION Sequence 201 from patent US 5693535.
ACCESSION I78403
VERSION I78403.1 GI:3014557
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 14)
AUTHORS Draper,K.G., Chowrira,B., McSwiggen,J., Stinchcomb,D.T. and
TITLE Thompson,J.D.
METHOD HIV targeted ribozymes
JOURNAL Patent: US 5693535-A 201 02-DEC-1997;
FEATURES Location/Qualifiers
source 1.14
BASE COUNT 7 a 3 c 3 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.1e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 13
LOCUS A14957 16 bp DNA PAT 18-FEB-1994
DEFINITION synthetic oligonucleotide (N2) from patent EP0211299.
ACCESSION A14957
VERSION A14957.1 GI:491869
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 16)
AUTHORS Habermann,P., Stengelin,S. and Wengenmayer,F.
TITLE Fusion proteins, method for their production and their use
JOURNAL Patent: EP 0211299-A 2 25-FEB-1987;
FEATURES Location/Qualifiers
source 1.16
BASE COUNT 1 a 4 c 2 g 9 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 16;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 15 CAATGAAGCAA 4

RESULT 14
LOCUS A14956 18 bp DNA PAT 18-FEB-1994
DEFINITION synthetic oligonucleotide (N1) from patent EP0211299.
ACCESSION A14956
VERSION A14956.1 GI:491868
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
REFERENCE 1 (bases 1 to 18)
AUTHORS Habermann,P., Stengelin,S. and Wengenmayer,F.
TITLE Fusion proteins, method for their production and their use
JOURNAL Patent: EP 0211299-A 1 25-FEB-1987;
FEATURES Location/Qualifiers
source 1.18
BASE COUNT 9 a 3 c 5 g 1 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagca 85
|||||
Db 4 CAATGAAGCAA 15

RESULT 15
LOCUS AR099365 18 bp DNA PAT 14-FEB-2001
DEFINITION Sequence 19 from patent US 6077709.

ACCESSION AR099365
VERSION AR099365.1 GI:12809131
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Bennett,C.Frank, Ackermann,E.J., Swayze,E.E. and Cowsett,L.M.
TITLE Antisense modulation of Survivin expression
JOURNAL Patent: US 6077709-A 19 20-JUN-2000;
FEATURES Location/Qualifiers
source 1..18
/organism="unknown"
BASE COUNT 1 a 3 c 3 g 11 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 51 tctattgttg 62
|||||
Db 1 TCTATTGTGG 12

RESULT 16
A07597 20 bp DNA PAT 09-JUL-1993
LOCUS A07597 Synthetic antisense oligonucleotide (5493-5512).
DEFINITION A07597
ACCESSION A07597.1 GI:413100
VERSION
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
REFERENCE 1 (bases 1 to 20)
ARTIFICIAL SEQUENCE.
AUTHORS Stropp,U., Baumgarten,J., Loebberding,A., Springer,W., Piel,N.,
Kretschmer,A., Koelbl,H. and Frommet,W.
TITLE Antisense-oligonucleotides for inhibiting the transactivator target
sequence (Tat) of HIV-1, and the synthesis of the transactivator protein
(Pat) of HIV-1, and their use
JOURNAL Patent: EP 0386563-A 4 12-SEP-1990;
JOURNAL BAYER AG
FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
BASE COUNT 8 a 4 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaa 85
|||||
Db 2 CAATGAAGCAA 13

RESULT 17
A64909/c 20 bp DNA PAT 29-MAR-1999
LOCUS A64909 Sequence 66 from Patent WO9731114.
DEFINITION A64909
ACCESSION A64909.1 GI:4530900
VERSION
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Burnham,M.K. and Hodgson,J.E.

TITLE POLYNUCLEOTIDES AND AMINOACID SEQUENCES FROM STAPHYLOCOCCUS AUREUS
JOURNAL Patent: WO 9731114-A 66 28-AUG-1997;
AUTHORS SMITHKLINE BEECHAM PLC (GB)
FEATURES Location/Qualifiers
source 1..20
/organism="unidentified"
BASE COUNT 11 a 4 c 2 g 3 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 20 GTTTCACGTTT 9

RESULT 18
AR043539 21 bp DNA PAT 29-SEP-1999
LOCUS AR043539 Sequence 6 from patent US 5814490.
DEFINITION AR043539
ACCESSION AR043539.1 GI:5964547
VERSION
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Spears,P.A.
TITLE Amplification and detection of chlamydia trachomatis nucleic acids
JOURNAL Patent: US 5814490-A 6 29-SEP-1998;
FEATURES Location/Qualifiers
source 1..21
/organism="unknown"
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 19
AR063857 21 bp DNA PAT 29-SEP-1999
LOCUS AR063857 Sequence 4 from patent US 5846726.
DEFINITION AR063857
ACCESSION AR063857.1 GI:5993165
VERSION
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Nadeau,J.G., Pither,J.Bruce, Schram,J.L., Linn,C.Preston, Vonk,G.P.
TITLE Detection of nucleic acids by fluorescence quenching
JOURNAL Patent: US 5846726-A 4 08-DEC-1998;
FEATURES Location/Qualifiers
source 1..21
/organism="unknown"
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 20
LOCUS AR075821/c 21 bp DNA PAT 30-AUG-2000

DEFINITION Sequence 4 from patent US 5958700.

ACCESSION AR075821

VERSION AR075821.1 GI:10002567

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Nadeau,J.G., Pitner,J.Bruce, Linn,C.Preston and Schram,J.L.

TITLE Detection of nucleic acids by fluorescence quenching

JOURNAL Patent: US 5958700-A 4 28-SEP-1999;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 21
LOCUS AR098733/c 21 bp DNA PAT 14-FEB-2001

DEFINITION Sequence 8 from patent US 6077669.

ACCESSION AR098733

VERSION AR098733.1 GI:12808499

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Little,M.C. and Vonk,G.P.

TITLE Kit and method for fluorescence based detection assay

JOURNAL Patent: US 6077669-A 8 20-JUN-2000;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 22
LOCUS AR112325/c 21 bp DNA PAT 16-MAY-2001

DEFINITION Sequence 5 from patent US 6130047.

ACCESSION AR112325

VERSION AR112325.1 GI:14092225

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 21)

AUTHORS Nadeau,J.G., Hsieh,H.V., Pitner,J.Bruce and Linn,C.Preston.

TITLE Detection of nucleic acids by fluorescence quenching

JOURNAL Patent: US 6130047-A 5 10-OCT-2000;

FEATURES Location/Qualifiers

source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 23
LOCUS E30454/c 21 bp DNA PAT 07-FEB-2001

DEFINITION Method for detecting target nucleic acid sequence and oligonucleotide.

ACCESSION E30454

VERSION E30454.1 GI:13025611

KEYWORDS JP 1999056380-A/4.

SOURCE unidentified.

ORGANISM unidentified.

REFERENCE 1 (bases 1 to 21)

AUTHORS James,G.N.J.J., Pitona,C.P.R.R. and L,S.

TITLE Method for detecting target nucleic acid sequence and

JOURNAL Patent: JP 1999056380-A 4 02-MAR-1999;

COMMENT BECTON DICKINSON & CO

OS Unidentified

PN JP 1999056380-A/4

PD 02-MAR-1999

PE 29-MAY-1998 JP 1998166141

PR 30-MAY-1997 US 08/865 675

PI JAMES G NADEAU,J BRUCE PITONA,C PRESTON RIN,JAMES L SHURAMU PC

CI2N15/09,CI2Q1/68,G01N33/50,G01N33/366,CI2N15/00 CC

Strandedness: Single;

CC Topology: Linear;

EH Key

FT source 1..21

BASE COUNT 8 a 5 c 3 g 5 t

ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 21;

Best Local Similarity 100.0%; Pred. No. 5.2e+04;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 39 gtttcattgttt 50
|||||
Db 13 GTTTCATGCTTT 2

RESULT 24
LOCUS E32364/c 21 bp DNA PAT 07-FEB-2001

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DEFINITION Method for detecting nucleic acid by fluorescent quenching.
ACCESSION E33634
VERSION E3364.1 GI:13026696
KEYWORDS JP 1999123083-A/4.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 21)
AUTHORS James,G.N.J., Pitona,J.L.S., Preston,R.G.P. and G.T.W.
TITLE Method for detecting nucleic acid by fluorescent quenching
JOURNAL Patent: JP 1999123083-A 4 11-MAY-1999;
BECTON DICKINSON & CO
COMMENT OS Unidentified
PN JP 1999123083-A/4
PD 11-MAY-1999
PF 13-MAY-1997 US 08/855 085
PI JAMES G NADEAU,J BLUCE PITONA,JAMES L SHURAMU,C PRESTON RIN,
PI GREN P VONG,
PI G TERANSU WALKER
PC C12N15/09,C07H21/00,C12Q1/68,G01N21/64,G01N33/58,C12N15/00 CC
Strandness: Single;
CC Topology: Linear;
FH Key Location/Qualifiers
FT source 1..21
/organism='Unidentified'.
FEATURES
SOURCE 1..21
Location/Qualifiers
/organism='unidentified'
/db_xref='taxon:32644'
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 25
E33635/c 21 bp DNA PAT 07-FEB-2001
DEFINITION Detection of nucleic acid by disappearance of fluorescence.
ACCESSION E33635
VERSION E33635.1 GI:13027030
KEYWORDS JP 1999155598-A/5.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 21)
AUTHORS James,G.N.H., C.J.B.P.P. and Preston,R.
TITLE Detection of nucleic acid by disappearance of fluorescence
JOURNAL Patent: JP 1999155598-A 5 15-JUN-1999;
BECTON DICKINSON & CO
COMMENT OS Artificial Sequence
PN JP 1999155598-A/5
PD 15-JUN-1999
PF 22-SEP-1998 JP 1998267492
PR 23-SEP-1997 US 08/933749
PI JAMES G NADEAU,HELEN V C,J BLUCE PITONA,C PRESTON RIN PC
C12Q1/68,C07H21/00,C12N15/09,G01N21/64,G01N33/542,G01N33/566, PC
C12N15/00
CC
FH Key Location/Qualifiers
FT source 1..21
Location/Qualifiers
/organism='Artificial Sequence'.
FEATURES
SOURCE 1..21
Location/Qualifiers
/organism='unidentified'

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BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 26
E35692/c 21 bp DNA PAT 07-FEB-2001
DEFINITION Detection assay with the use of fluorescence and kit therefor.
ACCESSION E35692
VERSION E35692.1 GI:13019164
KEYWORDS JP 1999225799-A/8.
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 21)
AUTHORS Michael,C.L.G.G. and Vong.
TITLE Detection assay with the use of fluorescence and kit therefor
JOURNAL Patent: JP 1999225799-A 8 24-AUG-1999;
BECTON DICKINSON & CO
COMMENT OS Artificial Sequence
PN JP 1999225799-A/8
PD 24-AUG-1999
PF 04-NOV-1998 JP 1998312790
PR 04-NOV-1997 US 08/964020
PI MICHAEL C LITTLE,GREN P VONG
PC C12Q1/68,G01N21/78,G01N33/50,G01N33/50,C12N15/09,C12N15/00 CC
FH Key Location/Qualifiers
FT source 1..21
Location/Qualifiers
/organism='Artificial Sequence'.
FEATURES
SOURCE 1..21
Location/Qualifiers
/organism='unidentified'
/db_xref='taxon:32644'
BASE COUNT 8 a 5 c 3 g 5 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 5.2e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 39 gtttcacgttt 50
|||||
Db 13 GTTTCACGTTT 2

RESULT 27
AR037890/c 23 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 10 from patent US 5804383.
ACCESSION AR037890
VERSION AR037890.1 GI:5956607
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 23)
AUTHORS Grenert,D.C. and Dohman,A.F.
TITLE Method and assay for detection of the expression of allele-specific
mutations by allele-specific in situ reverse transcriptase
polymerase chain reaction
JOURNAL Patent: US 5804383-A 10 08-SEP-1998;
FEATURES
SOURCE 1..21
Location/Qualifiers

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source 1. .23
/organism="unknown"
BASE COUNT 4 a 4 c 3 g 12 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 23;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 59 ttggagcaataa 70
|||||
Db 23 TTGGAGCAATAA 12

RESULT 28
AR054522 AR054522 24 bp DNA PAT 29-SEP-1999
LOCUS
DEFINITION Sequence 31 from patent US 5837441.
ACCESSION AR054522
VERSION AR054522.1 GI:5980099
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Hjelte, B. and Jensen, S.
TITLE Hantavirus-associated respiratory distress virus antigens
JOURNAL Patent: US 5837441-A 31 17-NOV-1998;
FEATURES
source 1. .24
/organism="unknown"

BASE COUNT 8 a 1 c 7 g 8 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 21 gtttactgtaac 32
|||||
Db 2 GTTTACTGTAAT 13

RESULT 29
AR151501 AR151501 24 bp DNA PAT 08-AUG-2001
LOCUS
DEFINITION Sequence 26 from patent US 6232094.
ACCESSION AR151501
VERSION AR151501.1 GI:15117551
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Hanson, L., Stromqvist, M., Bergstrom, S., Hernell, O., and Tornell, J.
TITLE DNA encoding human kappa casein and process for obtaining the protein
JOURNAL Patent: US 6232094-A 26 15-MAY-2001;
FEATURES
source 1. .24
/organism="unknown"
BASE COUNT 5 a 3 c 2 g 14 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 35 ttatgtttcat 46
|||||

Db 12 TTATGTTTTCAT 23

RESULT 30
AR082296/C AR082296 25 bp DNA PAT 31-AUG-2000
LOCUS
DEFINITION Sequence 140 from patent US 5972704.
ACCESSION AR082296
VERSION AR082296.1 GI:10009022
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowrira, B., McSwiggen, J., Stinchcomb, D.T. and
TITLE HIV net targeted ribozymes
JOURNAL Patent: US 5972704-A 140 26-OCT-1999;
FEATURES
source 1. .25
/organism="unknown"

BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaaa 85
|||||
Db 23 CAATGAAGCAA 12

RESULT 31
AR120838/C AR120838 25 bp DNA PAT 16-MAY-2001
LOCUS
DEFINITION Sequence 140 from patent US 6159692.
ACCESSION AR120838
VERSION AR120838.1 GI:14104414
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowrira, B., McSwiggen, J., Stinchcomb, D.T. and
TITLE Method and reagent for inhibiting human immunodeficiency virus replication
JOURNAL Patent: US 6159692-A 140 12-DEC-2000;
FEATURES
source 1. .25
/organism="unknown"

BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 74 caatgaagcaaa 85
|||||
Db 23 CAATGAAGCAA 12

RESULT 32
AX115268 AX115268 25 bp DNA PAT 11-MAY-2001
LOCUS
DEFINITION Sequence 391 from Patent WO0129262.
ACCESSION AX115268
VERSION AX115268.1 GI:14032210
KEYWORDS

SOURCE synthetic construct.
ORGANISM artificial sequence.
REFERENCE 1 (bases 1 to 25)
AUTHORS Picoult-Newburg, L. and Pohl, M.
TITLE Genotyping reagents, kits and methods of use thereof
JOURNAL Patent: WO 0129262-A 391 26-APR-2001;
Orchid Biosciences, Inc. (US)
FEATURES Location/Qualifiers
SOURCE 1..25
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Primer"
BASE COUNT 8 a 4 c 3 g 10 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 7 aaatattatca 18
|||||
Db 3 AAATTATTTC 14
RESULT 33
178342/c 178342 25 bp DNA PAT 03-APR-1998
LOCUS I78342 Sequence 140 from patent US 5693535.
DEFINITION I78342
ACCESSION I78342
VERSION I78342.1 GI:3014496
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Draper, K.G., Chowitra, B., McSwiggen, J., Stinchcomb, D.T. and
Thompson, J.D.
TITLE HIV targeted ribozymes
JOURNAL Patent: US 5693535-A 140 02-DEC-1997;
FEATURES Location/Qualifiers
SOURCE 1..25
/organism="unknown"
BASE COUNT 6 a 4 c 5 g 10 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 74 caatgaagca 85
|||||
Db 23 CAATGAAGCAA 12
RESULT 34
AR060384/c 27 bp DNA PAT 29-SEP-1999
LOCUS AR060384 Sequence 6 from patent US 5840568.
DEFINITION AR060384
ACCESSION AR060384
VERSION AR060384.1 GI:5986834
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfrendschuh, M.
TITLE Hodgkin's disease associated molecules and uses thereof
JOURNAL Patent: US 5840568-A 6 24-NOV-1998;
FEATURES Location/Qualifiers
SOURCE 1..27

BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88
|||||
Db 26 TGAAGCAAGTG 15
RESULT 35
AR117878/c 27 bp DNA PAT 16-MAY-2001
LOCUS AR117878 Sequence 6 from patent US 6140464.
DEFINITION AR117878
ACCESSION AR117878
VERSION AR117878.1 GI:14098784
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfrendschuh, M. and Rammensee, H.
TITLE Nonapeptides that bind a HLA-A2.1 molecule
JOURNAL Patent: US 6140464-A 6 31-OCT-2000;
FEATURES Location/Qualifiers
SOURCE 1..27
/organism="unknown"
BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88
|||||
Db 26 TGAAGCAAGTG 15
RESULT 36
AX114027/c 27 bp DNA PAT 08-MAY-2001
LOCUS AX114027 Sequence 6 from Patent EP1108432.
DEFINITION AX114027
ACCESSION AX114027
VERSION AX114027.1 GI:14018204
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 27)
AUTHORS Pfrendschuh, M.I.
TITLE Method for identifying or isolating a molecule and molecules
JOURNAL Patent: EP 1108432-A 6 20-JUN-2001;
JOURNAL identified thereby
LUDWIG INSTITUTE FOR CANCER RESEARCH (US)
FEATURES Location/Qualifiers
SOURCE 1..27
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 5 a 7 c 4 g 11 t
ORIGIN
Query Match 13.3%; Score 12; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 77 tgaagcaagt 88

Db 26 TGAAGCAAGTG 15

RESULT 37

AR082955 AR082955 28 bp DNA PAT 01-SEP-2000
 LOCUS Sequence 20 from patent US 5976795.
 DEFINITION AR082955
 ACCESSION AR082955
 VERSION AR082955.1 GI:10009745
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Voytas,D.F. and Zou,S.
 TITLE Retrotransposon and methods
 JOURNAL Patent: US 5976795-A 20 02-NOV-1999;
 FEATURES Location/Qualifiers
 source 1..28
 /organism="unknown"
 BASE COUNT 7 a 0 c 4 g 15 t 2 others
 ORIGIN

Query Match

Best Local Similarity 13.3%; Score 12; DB 6; Length 28;
 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 31 attttatgtt 42

Db 7 ATTTTATGTTT 18

RESULT 38
 AR082956/C AR082956 28 bp DNA PAT 01-SEP-2000
 LOCUS Sequence 21 from patent US 5976795.
 DEFINITION AR082956
 ACCESSION AR082956
 VERSION AR082956.1 GI:10009746
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 28)
 AUTHORS Voytas,D.F. and Zou,S.
 TITLE Retrotransposon and methods
 JOURNAL Patent: US 5976795-A 21 02-NOV-1999;
 FEATURES Location/Qualifiers
 source 1..28
 /organism="unknown"
 BASE COUNT 15 a 4 c 0 g 7 t 2 others
 ORIGIN

Query Match

Best Local Similarity 13.3%; Score 12; DB 6; Length 28;
 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 31 attttatgtt 42

Db 22 ATTTTATGTTT 11

RESULT 39
 AX155921/C AX155921 29 bp DNA PAT 22-JUN-2001
 LOCUS Sequence 164 from Patent WO0140474.
 DEFINITION AX155921
 ACCESSION AX155921
 VERSION AX155921.1 GI:14537028
 KEYWORDS
 SOURCE Chlamydia sp.
 ORGANISM Chlamydia sp.

REFERENCE 1 (bases 1 to 29)
 AUTHORS Probst,P., Bhatia,A., Skeiky,Y.A., Flinn,S.P. and Scholler,J.
 TITLE Compounds and methods for treatment and diagnosis of chlamydial
 JOURNAL Infection
 Patent: WO 0140474-A 164 07-JUN-2001;
 FEATURES CORIXA CORPORATION (US)
 source Location/Qualifiers
 1..29
 /organism="Chlamydia sp."
 /db_xref="taxon:35827"

BASE COUNT

12 a 4 c 4 g 9 t

ORIGIN

Query Match

Best Local Similarity 13.3%; Score 12; DB 6; Length 29;
 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 48 ttctctatgt 59

Db 15 TTTCTTATGTT 4

RESULT 40
 A43687/C A43687 30 bp DNA PAT 06-MAR-1997
 LOCUS Sequence 3 from Patent WO9508642.
 DEFINITION A43687
 ACCESSION A43687
 VERSION A43687.1 GI:2298875
 KEYWORDS
 SOURCE unidentified.
 ORGANISM unidentified.

REFERENCE 1 (bases 1 to 30)
 AUTHORS Sammes,P.G. and Garman,A.J.
 TITLE NUCLEIC ACID DETECTION WITH ENERGY TRANSFER
 JOURNAL Patent: WO 9508642-A 3 30-MAR-1995;
 ZENECA LTD (GB)

COMMENT Other publication AU 7662794 950410
 Other publication GB 2283095 950426.

FEATURES Location/Qualifiers
 source 1..30
 /organism="unidentified"
 /db_xref="taxon:32644"

BASE COUNT 12 a 4 c 4 g 10 t

ORIGIN

Query Match

Best Local Similarity 13.3%; Score 12; DB 6; Length 30;
 100.0%; Pred. No. 5.3e+04;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 26 ctgtaatttta 37

Db 29 CTGTATTTTTA 18

RESULT 41
 AR028176/C AR028176 30 bp DNA PAT 29-SEP-1999
 LOCUS Sequence 25 from patent US 5858661.
 DEFINITION AR028176
 ACCESSION AR028176
 VERSION AR028176.1 GI:5940149
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
 AUTHORS Shiloh,Y.
 TITLE Ataxia-telangiectasia gene and its genomic organization
 JOURNAL Patent: US 5858661-A 25 12-JAN-1999;
 Location/Qualifiers

FEATURES

source 1..30
/organism="unknown"
BASE COUNT 8 a 2 c 2 g 18 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
|||||
DB 19 AAAAAATTATT 8

RESULT 42
AR050260/c 30 bp DNA PAT 29-SEP-1999
LOCUS AR050260 Sequence 3 from patent US 5827653.
DEFINITION AR050260
ACCESSION AR050260
VERSION AR050260.1 GI:5972985
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Sammes,P.George and Garman,A.John.
TITLE Nucleic acid detection with energy transfer
JOURNAL Patent: US 5827653-A 3 27-OCT-1998;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 12 a 4 c 4 g 10 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ctgtaattttta 37
|||||
DB 29 CTGTAATTTTAA 18

RESULT 43
ARI38579/c 30 bp DNA PAT 16-JUN-2001
LOCUS ARI38579 Sequence 104 from patent US 6200749.
DEFINITION ARI38579
ACCESSION ARI38579
VERSION ARI38579.1 GI:14480924
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Shiloh,Y.
TITLE Mutated forms of the ataxia-telangiectasia gene and method to
screen for a partial A-T phenotype
JOURNAL Patent: US 6200749-A 104 13-MAR-2001;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 8 a 2 c 2 g 18 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 aaaaaattatt 15
|||||

DB 19 AAAAAATTATT 8

RESULT 44
ARI40326/c 30 bp DNA PAT 16-JUN-2001
LOCUS ARI40326 Sequence 14 from patent US 6207455.
DEFINITION ARI40326
ACCESSION ARI40326
VERSION ARI40326.1 GI:14482822
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 30)
AUTHORS Chang,L.-J.
TITLE Lenticviral vectors
JOURNAL Patent: US 6207455-A 14 27-MAR-2001;
FEATURES Location/Qualifiers
source 1..30
/organism="unknown"

BASE COUNT 7 a 5 c 6 g 12 t
ORIGIN

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 caatgaagcaaa 85
|||||
DB 28 CAATGAAGCAAA 17

RESULT 45
AX020976/c 30 bp DNA PAT 07-SEP-2000
LOCUS AX020976 Sequence 22 from Patent EP0928832.
DEFINITION AX020976
ACCESSION AX020976
VERSION AX020976.1 GI:10044639
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequence.

REFERENCE 1 (bases 1 to 30)
AUTHORS Kelly,S.J., Weston,S.L. and Robertson,N.H.
TITLE Cyclic fibrosis test based on the detection of mutations in the
cife gene by arms
JOURNAL Patent: EP 0928832-A 22 14-JUL-1999;
FEATURES ZENECA LTD (GB)
source 1..30
Location/Qualifiers

BASE COUNT 8 a 6 c 5 g 11 t
ORIGIN
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="PCR Amplification Primer"

Query Match 13.3%; Score 12; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 5.3e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 59 ttgaggaataa 70
|||||
DB 17 TTGAGACATAA 6

Search completed: January 24, 2002, 03:22:38
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BASE COUNT 2 a 5 c 3 g 14 t
 ORIGIN /db_xref="taxon:32630"
 /note="synthetic oligonucleotide"

Query Match 14.4%; Score 13; DB 6; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 34 ttatgtttcat 46
 Db 8 tttatgtttcat 20

RESULT 2
 AX039624 26 bp DNA PAT 18-NOV-2000
 LOCUS AX039624
 DEFINITION Sequence 13 from Patent WO0063441.
 ACCESSION AX039624
 VERSION AX039624.1 GI:11229653
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM artificial sequence.
 REFERENCE 1 (bases 1 to 26)
 AUTHORS Herinstad,C. and Davis,R.E.
 TITLE Single nucleotide polymorphisms in mitochondrial genes that segreg
 JOURNAL Patent: WO 0063441-A 13 26-OCT-2000;
 MITOKOR (US)

FEATURES
 source Location/Qualifiers
 1..26
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="PCR primer"

BASE COUNT 8 a 2 c 5 g 11 t
 ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 37 atgtttcatgtt 49
 Db 7 atgtttcatgtt 19

RESULT 3
 AX039654 26 bp DNA PAT 18-NOV-2000
 LOCUS AX039654
 DEFINITION Sequence 43 from Patent WO0063441.
 ACCESSION AX039654
 VERSION AX039654.1 GI:11229663
 KEYWORDS
 SOURCE synthetic construct.
 ORGANISM artificial sequence.
 REFERENCE 1 (bases 1 to 26)
 AUTHORS Herinstad,C. and Davis,R.E.
 TITLE Single nucleotide polymorphisms in mitochondrial genes that segreg
 JOURNAL Patent: WO 0063441-A 43 26-OCT-2000;
 MITOKOR (US)

FEATURES
 source Location/Qualifiers
 1..26
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 /note="PCR primer"

BASE COUNT 8 a 2 c 5 g 11 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 26;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 atgtttcatgtt 49
 Db 7 atgtttcatgtt 19

RESULT 4
 AR028230 30 bp DNA PAT 29-SEP-1999
 LOCUS AR028230
 DEFINITION Sequence 79 from patent US 5858661.
 ACCESSION AR028230
 VERSION AR028230.1 GI:5940203
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 30)
 AUTHORS Shiloh,Y.
 TITLE Ataxia-telangiectasia gene and its genomic organization
 JOURNAL Patent: US 5858661-A 79 12-JAN-1999;
 FEATURES Location/Qualifiers
 source 1..30
 /organism="unknown"

BASE COUNT 13 a 2 c 1 g 14 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 5 aaaaattatc 17
 Db 15 AAAAAATTATTC 27

RESULT 5
 ARI38633 30 bp DNA PAT 16-JUN-2001
 LOCUS ARI38633
 DEFINITION Sequence 158 from patent US 6200749.
 ACCESSION ARI38633
 VERSION ARI38633.1 GI:14480978
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 30)
 AUTHORS Shiloh,Y.
 TITLE Mutated forms of the ataxia-telangiectasia gene and method to
 screen for a partial A-T phenotype
 JOURNAL Patent: US 6200749-A 158 13-MAR-2001;
 FEATURES Location/Qualifiers
 source 1..30
 /organism="unknown"

BASE COUNT 13 a 2 c 1 g 14 t
 ORIGIN

Query Match 14.4%; Score 13; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 aaaaattatc 17
 Db 15 AAAAAATTATTC 27

RESULT 6
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us-09-531-438-3.oli.rng

OM nucleic - nucleic search, using sw model

Run on: January 24, 2002, 02:27:08 ; Search time 222.28 Seconds

(without alignments)
1261.226 Million cell updates/sec

Title: US-09-531-438-3

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Gapop 60.0 , Gapext 60.0

Searched: 930621.seqs, 428662619 residues

Word size :

Total number of hits satisfying chosen parameters:	989696
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Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Listing first 45 summaries

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Pred. No.' is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	16	4.9	29	17	AA74255	Primer for amplif
2	16	4.9	29	21	AA51200	N-terminal primer
3	16	4.9	32	20	AA227689	PCR primer for Ver
4	16	4.9	36	18	AA797603	<i>Shigella dysenteriae</i> Primer Y104F. Syn
5	16	4.9	39	20	AAV90775	Human map-related
6	16	4.9	47	21	AA268379	Dog genomic marker
7	15	4.6	20	21	AAA66183	Human high motilit
8	15	4.6	24	22	AA446804	Hel-N2 selected se
9	15	4.6	27	17	AAT39439	Human Hel-N2 selec
10	15	4.6	27	19	AAV37457	Triple helix formi
11	15	4.6	30	14	AAQ43975	

12	15	4.6	48.16	AAT25576	Human gene signatu
13	14	4.3	18.13	AAQ20161	Cross-linking olig
14	14	4.3	18.13	AAQ20160	Cross-linking olig
15	14	4.3	18.13	AAQ30311	Oligomer HSV724 fo
16	14	4.3	18.13	AAQ30310	Oligomer HSV723 fo
17	14	4.3	18.20	AAZ22162	Human c-IAP-1 mRNA
18	14	4.3	22.22	AAAS01590	Human IQGAP2 Cpg 1
19	14	4.3	22.22	AAAS01643	Human IQGAP2 5'-UTR
20	14	4.3	26.19	AAV07952	Helicobacter pylori
21	14	4.3	26.19	AAV07952	Helicobacter pylori
22	14	4.3	27.19	AAV07937	Helicobacter pylori
23	14	4.3	29.17	AAV12655	Primer for amplify
24	14	4.3	29.21	AAAS12000	N-terminal primer
25	14	4.3	31.16	AAAT35703	Human gene signatu
26	14	4.3	31.19	AAV67854	Nucleotide fragmen
27	14	4.3	32.20	AAZ27689	PCR primer for Ver
28	14	4.3	36.18	AAAT97603	Shigella dysenter
29	14	4.3	36.22	AAAC90606	Tomato spotted wil
30	14	4.3	37.15	AAAG62952	Oligonucleotide us
31	14	4.3	45.22	AAAF55449	Oligonucleotide us
32	14	4.3	45.22	AAAF55450	Oligonucleotide us
33	14	4.3	45.22	AAAC88874	Oligonucleotide TA
34	14	4.3	45.22	AAAC88875	Oligonucleotide TA
35	14	4.3	47.21	AAZ66366	Human map-related
36	14	4.3	47.21	AAZ67473	Human map-related
37	14	4.3	47.21	AAZ67533	Human map-related
38	14	4.3	47.21	AAZ67549	Human map-related
39	14	4.3	47.21	AAZ67813	Human map-related
40	14	4.3	50.21	AAAF98312	Human MSH6 fragmen
41	13	4.0	15.22	AAAF48097	IGFBP3 oligonucleo
42	13	4.0	15.22	AAAF48098	IGFBP3 oligonucleo
43	13	4.0	15.22	AAAF48099	IGFBP3 oligonucleo
44	13	4.0	16.21	AAAF57758	Nucleotide sequenc
45	13	4.0	17.16	AAQ920084	Renilla reniformis

ALIGNMENTS

RESULT	1
AAT42655	
ID AAT42655	standard; DNA; 29 BP.
XX AC	
AC	
AAT42655;	
DDT	
XX	
XX	
XX	
DE	
XX	
XX	
KW	
XX	
XX	
XX	
S	
SS	
NN	
CN	
XX	
XX	
XX	
PD	

Pred. No.' is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

PS Example 6; Page 58; 101pp; English.

XX Compositions containing neutralizing antitoxin against one or more E.
CC coli verotoxin (VT) can be used to treat intoxicated adults and
CC children with enteric bacterial infections. They may also be used as
CC prophylactics e.g. as a vaccine, against diarrhoeal disease or the
CC development of extra-intestinal complications of E.coli infection,
CC especially haemolytic uremic syndrome. The antitoxin can also be
CC used to detect E. coli VT in a sample. The VT is recombinant,
CC preferably a fusion protein containing a non-VT protein sequence and
CC part of the E.coli VT1 or VT2 sequence. Two primers (AA142655,
CC AA142656) were used to amplify the verotoxin VT-1 A subunit coding
CC sequence and add a histidine tag coding sequence to the subunit
CC sequence. Two primers (AA142655, AA142658) were used to amplify the
CC verotoxin VT-1 A and B subunits and add a histidine tag coding
CC sequence to the subunit sequences.

XX Sequence 29 BP: 11 A; 2 C; 5 G; 11 T; 0 other:

Query Match 4.9%; Score 16; DB 17; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204

Db 9 aaataattatttta 24

RESULT 2

AA27689 standard; DNA: 29 BP.

AC AA27689;

DT 26-SEP-2000 (first entry)

DE N-terminal primer for E. coli verotoxin 1 subunit A gene.

XX VT-1; verotoxin; antitoxin therapy; fusion protein; affinity tag; food;

KM recombinant production; screening; dairy; anti-bacterial; vaccine;

KM primer: polyhistidine; ss.

XX Escherichia coli.

OS Synthetic.

PN US6080400-A.

XX 27-JUN-2000.

PF 13-MAR-1997; 97US-0816977.

XX 24-MAR-1995; 95US-0410058.

XX (OPHT-) OPHIDIAN PHARM INC.

PA Williams JA, Byrne LM;

PI WPI: 2000-451195/39.

XX Bacterial cell for recombinantly expressing bacterial toxins in large
PT quantities useful for immunization and treatment of bacterial
PT infections, comprises expression vector encoding bacterial toxin

PS Example 6; Column 83; 83pp; English.

XX E. coli verotoxin (VT) type 1 and 2 subunits A and B were cloned into
CC PET-23b, designed to allow expression of the native proteins containing
C-terminal polyhistidine tags. The VT-1 and VT-2 genes were engineered
to convert the signal sequence methionine codon into a NdeI site to
allow cloning of the amplified genes into the vector without addition of
an encoded amino acids. The C-terminal primers comprises the
17 codons of each gene fused to the sequence CTCGACC, in order

to add the polyhistidine tag. The primers delete the native stop codons,
CC and when cloned into PET-23 add a C-terminal extension of Leu-Glu-(His)₆.
CC VT-B chains are small proteins (approximately 8 kDa), so use of a small
CC affinity tag was preferred (i.e. polyhistidine). A polyhistidine affinity
CC tag facilitates single-step affinity purification of subunits from
CC periplasmic extracts. However, due to poor recovery of his-tagged VT-1 A
CC and VT-2 A chains, expression of maltose binding protein (MBP) fused
CC subunits was undertaken. Due to the toxicity of the VT-2 B subunit,
CC strict uninduced promoter control is necessary to permit cell viability.
CC Bacterial host cells expressing a recombinant expression vector encoding
CC a polyhistidine affinity tag and a portion of the VT-2 B chain are
CC claimed. The vector is chosen from PET24hisVT2BL+, PET24hisVT2BL- and
CC PET24hisVT2B, where "L+" indicates that the vector encodes the preprotein
CC form of the protein and "L-" indicates that the vector encodes the mature
CC form of the protein. The bacterial cell is capable of expressing large
CC quantities (40 mg/l) of VT-2B. The toxins are useful for immunizing
CC non-mammals and for detecting bacterial toxins in environmental samples
CC including soil, water, industrial samples, biological samples and samples
CC obtained from food and dairy processing instruments.

XX Sequence 29 BP: 11 A; 2 C; 5 G; 11 T; 0 other:

Query Match 4.9%; Score 16; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1e+03;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 189 aaataattatttta 204

Db 9 aaataattatttta 24

RESULT 3

AA27689 standard; DNA: 32 BP.

AC AA27689;

DT 22-DEC-1999 (first entry)

DE PCR primer for Verotoxin gene.

XX Verotoxin; VT1; VT2; detection; PCR primer; ss.

XX Escherichia coli.

OS Synthetic.

PN JP11243996-A.

XX 14-SEP-1999.

PF 27-FEB-1998; 98JP-0047677.

XX 27-FEB-1998; 98JP-0047677.

XX (TOYM) TOYBO KK.

PA WPI: 1999-603716/52.

XX An oligonucleotide for amplification of verotoxin - useful in the
PT detection of inactivated verotoxin gene by transfer of a foreign DNA
PT fragment

PS Claim 11; Page 9; 10pp; Japanese.

XX This sequence represents a PCR primer of the invention. The primer is
CC used for amplification of the E. coli verotoxin (VT) gene. The
CC oligonucleotide is useful for detection of inactivated VT gene by
CC transfer of a foreign DNA fragment. Simple, rapid and specific
CC amplification of VT gene from environmental factors is achieved using the
CC oligonucleotide of the invention.

XX Sequence 32 BP: 12 A; 2 C; 4 G; 14 T; 0 other:

Query Match 4.38; Score 15; DB 15; Length 48;
 Best Local Similarity 100.0%; Pred. No. 2.5e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 159 aaaaataataaaaa 173
 Db 17 aaaaataataaaaa 31

RESULT 13
 AAQ20161
 ID AAQ20161 standard; DNA; 18 BP.
 XX AC AAQ20161;
 XX XX
 DT 01-APR-1992 (first entry)
 XX XX
 DE Cross-linking oligomer 724 to target Herpes Simplex Virus 1.
 XX XX
 KW deoxyribonucleic acid; major groove; HSV;
 KW inverted polarity region; covalent cross-linking group; ss.
 XX OS Synthetic.
 XX FH
 FT Key
 FT modified_base 1
 FT Location/Qualifiers
 FT /tag= a
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 2
 FT /tag= b
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 3
 FT /tag= c
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 4
 FT /tag= d
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 5
 FT /tag= e
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 6
 FT /tag= f
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 7
 FT /tag= g
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 8
 FT /tag= h
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 9
 FT /tag= i
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 10
 FT /tag= j
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT misc_feature 11
 FT /tag= k
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 12
 FT /tag= l
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 13
 FT /tag= m
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"

FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT 15
 FT /tag= m
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT 17
 FT /tag= n
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT XX
 PN W09118997-A.
 XX XX
 PD 12-DEC-1991.
 XX XX
 PF 24-MAY-1991; 91WO-1003680.
 XX XX
 PR 14-JAN-1991; 91US-0640654.
 XX XX
 PR 25-MAY-1990; 90US-0529346.
 XX XX
 PA (GILE-) GILEAD SCIE INC.
 XX XX
 PI Matteucci MD, Krawczyk S;
 XX XX
 DR WPI; 1992-007480/01.
 XX XX
 PT New sequence-specific non-photo-activated crosslinking agents -
 PT bind to the major groove of duplex DNA and are esp. useful for
 PT treating latent infections e.g. HIV
 XX XX
 PS Example 4; Page 29; 42pp; English.
 XX XX
 CC This oligomer contains an inverted polarity region formed from an
 CC o-xylosa dimer synthon. Residues 11 and 12 are linked via an
 CC o-xylosa group (i.e. nucleotides that have xylose sugar linked via
 CC the o-xylosa ring). The sequence is designed to target the Herpes
 CC Simplex virus 1 beginning at nucleotide 10996 and to covalently
 CC cross-link to it. See also AAQ20151-Q20160.
 XX XX
 SO Sequence 18 BP; 12 A; 1 C; 0 G; 5 T; 0 other;

Query Match 4.38; Score 14; DB 13; Length 18;
 Best Local Similarity 100.0%; Pred. No. 6.9e+03;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 159 aaaaataataaaaa 172
 Db 2 aaaaataataaaaa 15

RESULT 14
 AAQ20160
 ID AAQ20160 standard; DNA; 18 BP.
 XX AC AAQ20160;
 XX XX
 DT 01-APR-1992 (first entry)
 XX XX
 DE Cross-linking oligomer 723 to target Herpes Simplex Virus 1.
 XX XX
 KW deoxyribonucleic acid; major groove; HSV;
 KW inverted polarity region; covalent cross-linking group; ss.
 XX OS Synthetic.
 XX FH
 FT Key
 FT modified_base 1
 FT Location/Qualifiers
 FT /tag= a
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"
 FT modified_base 2
 FT /tag= b
 FT /mod_base= OTHER
 FT /note= "N-methyl-8-oxo-2'-deoxyadenine"